PROTOCADHERIN MATERIALS AND METHODS

This application is a continuation-in-part of International Patent Application No. PCT/US93/12588 filed December 23, 1993 which is in turn a continuation-in-part of U.S. Patent Application Serial No. 07/998,003 which was filed on December 29, 1992.

FIELD OF THE INVENTION

The present invention relates, in general, to materials and methods relevant to cell-cell adhesion. More particularly, the invention relates to novel adhesion proteins, designated protocadherins, and to polynucleotide sequences encoding the protocadherins. The invention also relates to methods for inhibiting binding of the protocadherins to their natural ligands/antiligands.

BACKGROUND

In vivo, intercellular adhesion plays an important role in a wide range of events including morphogenesis and organ formation, leukocyte extravasion, tumor metastasis and invasion, and the formation of cell junctions. Additionally, cell-cell adhesion is crucial for the maintenance of tissue integrity.

Intercellular adhesion is mediated by specific cell surface adhesion molecules. Cell adhesion molecules have been classified into at least four families including the immunoglobulin superfamily, the integrin superfamily, the selectin family and the cadherin superfamily. All cell types that form solid tissues express some members of the cadherin superfamily suggesting that cadherins are involved in selective adhesion of most cell types.

Cadherins have been generally described as glycosylated integral membrane proteins that have an N-terminal extracellular domain (the N-terminal 113 amino acids of the domain appear to be directly involved in binding) consisting of five subdomains characterized by sequences unique to cadherins, a hydrophobic membrane-spanning domain and a C-terminal cytoplasmic domain that interacts with the cytoskeleton through catenins and other cytoskeleton-

10

5

15

20

associated proteins. Some cadherins lack a cytoplasmic domain, however, and appear to function in cell-cell adhesion by a different mechanism than cadherins having a cytoplasmic domain. The cytoplasmic domain is required for the adhesive function of the extracellular domain in cadherins that do have an cytoplasmic domain. Binding between members of the cadherin family expressed on different cells is homophilic (i.e., a member of the cadherin family binds to cadherins of its own or a closely related subclass) and Ca²⁺-dependent. For recent reviews on cadherins, see Takeichi, Annu. Rev. Biochem., 59: 237-252 (1990) and Takeichi, Science, 251: 1451-1455 (1991).

10

5

The first cadherins to be described (E-cadherin in mouse epithelial cells, L-CAM in avian liver, uvomorulin in the mouse blastocyst, and CAM 120/80 in human epithelial cells) were identified by their involvement in Ca²⁺-dependent cell adhesion and their unique immunological characteristics and tissue localization. With the later immunological identification of N-cadherin, which was found to have a different tissue distribution than E-cadherin, it became apparent that a new family of Ca²⁺-dependent cell-cell adhesion molecules had been discovered.

20

25

15

The molecular cloning of the genes encoding E-cadherin [see Nagafuchi et al., Nature, 329: 341-343 (1987)], N-cadherin [Hatta et al., J. Cell. Biol., 106: 873-881 (1988)], and P-cadherin [Nose et al., EMBO J., 6: 3655-3661 (1987)] provided structural evidence that the cadherins comprised a family of cell adhesion molecules. Cloning of L-CAM [Gallin et al., Proc. Natl. Acad., Sci. USA, 84: 2808-2812 (1987)] and uvomorulin [Ringwald et al., EMBO J., 6: 3647-3653 (1986)] revealed that they were identical to E-cadherin. Comparisons of the amino acid sequences of E-, N-, and P-cadherins showed a level of amino acid similarity of about 45%-58% among the three subclasses. Liaw et al., EMBO J., 9: 2701-2708 (1990) describes the use of PCR with degenerate oligonucleotides based on conserved regions of the E-, N- and P-cadherins to amplify N- and P-cadherin from a bovine microvascular endothelial cell cDNA.

The isolation by PCR of eight additional cadherins was reported in Suzuki et al., Cell Regulation, 2: 261-270 (1991). Subsequently, several other cadherins were described including R-cadherin [Inuzuka et al., Neuron, 7: 69-79 (1991)], M-cadherin [Donalies, Proc. Natl. Acad. Sci. USA, 88: 8024-8028 (1991)], B-cadherin [Napolitano, J. Cell. Biol., 113: 893-905 (1991)] and T-cadherin [Ranscht, Neuron, 7: 391-402 (1991)].

Additionally, proteins distantly related to cadherins such as desmoglein [Goodwin et al., Biochem. Biophys. Res. Commun., 173: 1224-1230 (1990) and Koch et al., Eur. J. Cell Biol., 53: 1-12 (1990)] and the desmocollins [Holton et al., J. Cell Science, 97: 239-246 (1990)] have been described. The extracellular domains of these molecules are structurally related to the extracellular domains of typical cadherins, but each has a unique cytoplasmic domain. Mahoney et al., Cell, 67: 853-868 (1991) describes a tumor suppressor gene of Drosophila, called fat, that also encodes a cadherin-related protein. The fat tumor suppressor comprises 34 cadherin-like subdomains followed by four EGF-like repeats, a transmembrane domain, and a novel cytoplasmic domain. The identification of these cadherin-related proteins is evidence that a large superfamily characterized by a cadherin extracellular domain motif exists.

Studies of the tissue expression of the various cadherin-related proteins reveal that each subclass of molecule has a unique tissue distribution pattern. For example, E-cadherin is found in epithelial cells while N-cadherin is found in neural and muscle cells. Expression of cadherin-related proteins also appears to be spatially and temporally regulated during development because individual proteins appear to be expressed by specific cells and tissues at specific developmental stages [for review see Takeichi (1991), supra]. Both the ectopic expression of cadherin-related proteins and the inhibition of native expression of cadherin-related proteins hinders the formation of normal tissue structure [Detrick et al., Neuron, 4: 493-506 (1990); Fujimori et al., Development, 110: 97-104 (1990); Kintner, Cell, 69: 225-236 (1992)].

10.

5

15

25

10

15

20

25

The unique temporal and tissue expression pattern of the different cadherins and cadherin-related proteins is particularly significant when the role each subclass of proteins may play in vivo in normal events (e.g., the maintenance of the intestinal epithelial barrier) and in abnormal events (e.g., tumor metastasis or inflammation) is considered. Different subclasses or combinations of subclasses of cadherin-related proteins are likely to be responsible for different cell-cell adhesion events in which therapeutic detection and/or intervention may For example, auto-antibodies from patients with pemphigus be desirable. vulgaris, an autoimmune skin disease characterized by blister formation caused by loss of cell adhesion, react with a cadherin-related protein offering direct support for adhesion function of cadherins in vivo [Amagai et al., Cell, 67: 869-877 (1991)]. Studies have also suggested that cadherins and cadherin-related proteins may have regulatory functions in addition to adhesive activity. Matsunaga et al., Nature, 334: 62-64 (1988) reports that N-cadherin has neurite outgrowth promoting activity. The Drosophila fat tumor supressor gene appears to regulate cell growth and supress tumor invasion as does mammalian E-cadherin [see Mahoney et al., supra; Frixen et al., J. Cell. Biol., 113:173-185 (1991); Chen et al., J. Cell, Biol., 114:319-327 (1991); and Vleminckx et al., Cell, 66:107-119 (1991)]. Thus, therapeutic intervention in the regulatory activities of cadherin-related proteins expressed in specific tissues may be desirable.

There thus continues to exist a need in the art for the identification and characterization of additional cadherin-related proteins which participate in cell-cell adhesion and/or regulatory events. Moreover, to the extent that cadherin-related proteins might form the basis for the development of therapeutic and diagnostic agents, it is essential that the genes encoding the proteins be cloned. Information about the DNA sequences and amino acid sequences encoding the cadherin-related proteins would provide for the large scale production of the proteins by recombinant techniques and for the identification of the tissues/cells naturally producing the proteins. Such sequence information would also permit

the preparation of antibody substances or other novel binding molecules specifically reactive with the cadherin-related proteins that may be useful in modulating the natural ligand/antiligand binding reactions in which the proteins are involved.

5

SUMMARY OF THE INVENTION

methods that are relevant to cell-cell adhesion. In one of its aspects, the present

invention provides purified and isolated polynucleotides (e.g., DNA and RNA, both sense and antisense strands) encoding the novel cell adhesion molecules

designated herein as protocadherins, including protocadherin-42, protocadherin-

The present invention provides cadherin-related materials and

10

15

20

25

43, protocadherin pc3, protocadherin pc4 and protocadherin pc5. Preferred polynucleotide sequences of the invention include genomic and cDNA sequences as well as wholly or partially synthesized DNA sequences, and biological replicas thereof (i.e., copies of the sequences made in vitro). Biologically active vectors comprising the polynucleotide sequences are also contemplated.

Specifically illustrating protocadherin polynucleotide sequences of the present invention are the inserts in the plasmids pRC/RSV-pc42 and pRC/RSV-pc43 which were deposited with the American Type Culture Collection

(ATCC), 12301 Parklawn Drive, Rockville, Maryland 20852 on December 16,

1992 and were assigned ATCC Accession Nos. 69162 and 69163, respectively.

The scientific value of the information contributed through the disclosures of the DNA and amino acid sequences of the present invention is manifest. For example, knowledge of the sequence of a partial or complete DNA encoding a protocadherin makes possible the isolation by standard DNA/DNA hybridization or PCR techniques of full length cDNA or genomic DNA sequences that encode the protein (or variants thereof) and, in the case of genomic DNA sequences, that specify protocadherin-specific regulatory sequences such as promoters, enhancers and the like. Alternatively, DNA sequences of the present invention may be chemically synthesized by conventional techniques.

10

15

Hybridization and PCR techiques also allow the isolation of DNAs encoding heterologous species proteins homologous to the protocadherins specifically illustrated herein.

According to another aspect of the invention, host cells, especially eucaryotic and procaryotic cells, are stably transformed or transfected with the polynucleotide sequences of the invention in a manner allowing the expression of protocadherin polypeptides in the cells. Host cells expressing protocadherin polypeptide products, when grown in a suitable culture medium, are particularly useful for the large scale production of protocadherin polypeptides, fragments and variants thereby enabling the isolation of the desired polypeptide products from the cells or from the medium in which the cells are grown.

The novel protocadherin protein products of the invention may be obtained as isolates from natural tissue sources, but are preferably produced by recombinant procedures involving the host cells of the invention. The products may be obtained in fully or partially glycosylated, partially or wholly deglycosylated, or non-glycosylated forms depending on the host cell selected or recombinant production and/or post-isolation processing.

Protocadherin variants according to the invention may comprise polypeptide analogs wherein one or more of the specified amino acids is deleted or replaced or wherein one or more non-naturally encoded amino acids are added: (1) without loss, and preferably with enhancement, of one or more of the biological activities or immunological characteristics specific for a protocadherin; or (2) with specific disablement of a particular ligand/antiligand binding function. Also contemplated by the present invention are antibody substances (e.g., monoclonal and polyclonal antibodies, chimeric and humanized antibodies, antibody domains including Fab, Fab', F(ab')₂, Fv or single variable domains, and single chain antibodies) which are specific for the protocadherins of the invention. Antibody substances can be developed using isolated natural, recombinant or synthetic protocadherin polypeptide products or host cells

20

expressing such products on their surfaces. The antibody substances may be utilized for purifying protocadherin polypeptides of the invention, for determining tissue expression of polypeptides and as antagonists of the ligand/antiligand binding activities of the protocadherins. Specifically illustrating monoclonal antibodies of the present invention are the protocadherin-43 specific monoclonal antibodies produced by the hybridoma cell line designated 38I2C which was deposited with the ATCC on December 2, 1992 and was assigned ATCC Accession No. HB 11207.

Numerous other aspects and advantages of the present invention will be apparent upon consideration of the following detailed description, reference being made to the drawing wherein FIGURE 1A-C is an alignment of protocadherin amino acid sequences of the invention with the amino acid sequences of N-cadherin and of the *Drosophila fat* tumor suppressor.

DETAILED DESCRIPTION

15

20

25

5

10

The present invention is illustrated by the following examples wherein Examples 1, 2 and 3 describe the isolation by PCR of protocadherin polynucleotide sequences. Example 3 also describes the chromosome localization of several protocadherin genes of the invention. Example 4 describes the isolation by DNA/DNA hybridization of additional protocadherin polynucleotide sequences of the present invention. Example 5 presents the construction of expression plasmids including polynucleotides encoding protocadherin-42 or protocadherin-43 and the transfection of L cells with the plasmids. The generation of antibodies to protocadherin-42 and protocadherin-43 is described in Example 6. Example 7 presents the results of immunoassays of transfected L cells for the expression of protocadherin-42 or protocadherin-43. Example 8 describes the cell aggregation properties of L cells transfected with protocadherin-42, protocadherin-43 or a chimeric protocadherin-43/E-cadherin molecule. The calcium-binding properties of pc43 are described in Example 9. The results of assays of various tissues and cell lines for the expression of protocadherin-42 and protocadherin-43

by Northern blot, Western blot and *in situ* hybridization are respectively presented in Examples 10, 11 and 12. Example 13 describes immunoprecipitation experiments identifying a 120 kDa protein that coprecipitates with protocadherin-43.

5

Example 1

The polymerase chain reaction (PCR) was used to isolate novel rat cDNA fragments encoding cadherin-related polypeptides.

Design of PCR Primers

Two regions of conserved amino acid sequence, one from the middle of the third cadherin extracellular subdomain (EC-3) and the other from the C-terminus of the fourth extracellular subdomain (EC-4), were identified by comparison of the published amino acid sequences for L-CAM (Gallin et al., supra), E-cadherin (Nagafuchi et al., supra), mouse P-cadherin (Nose et al., supra), uvomorulin (Ringwald et al., supra), chicken N-cadherin (Hatta et al., supra), mouse N-cadherin [Miyatani et al., Science, 245:631-635 (1989)] and human P-cadherin [Shimoyama et al., J. Cell. Biol., 109:1787-1794 (1989)], and the corresponding degenerate oligonucleotides respectively set out below in IUPAC-IUB Biochemical nomenclature were designed for use as PCR primers.

Primer 1 (SEO ID NO: 1)

5' AARSSNNTNGAYTRYGA 3'

Primer 2 (SEQ ID NO: 2)

3' TTRCTRTTRCGNGGNNN 5'

The degenerate oligonucleotides were synthesized using an Applied Biosystems model 380B DNA synthesizer (Foster City, California).

25 Cloning of cDNA Sequences by PCR

PCR was carried out in a manner similar to that described in Suzuki et al., Cell Regulation, 2: 261-270 (1991) on a rat brain cDNA preparation. Total RNA was prepared from rat brain by the guanidium

10

15

10 -

15

20

25

isothiocyanate/cesium chloride method described in Maniatis et al., pp. 196 in Molecular Cloning: A Laboratory Manual, Cold Spring Harbor, New York: Cold Spring Harbor Laboratory (1982). Brain poly(A)+ RNAs were then isolated using a FastTrack® kit (Invitrogen, San Diego, California) and cDNA was prepared using a cDNA synthesis kit (Boehringer Mannheim Biochemicals, Indianapolis, Indiana). The PCR reaction was initiated by adding 2.5 units of Taq DNA polymerase (Boehringer Mannheim Biochemicals) to 100 ng template cDNA and 10 μg of each primer, after which 35 reaction cycles of denaturation at 94°C for 1.5 minutes, annealing at 45°C for 2 minutes, and polymerization at 72°C for 3 minutes were carried out. Two major bands of about 450 base pairs (bp) and 130 bp in size were found when the products of the PCR reaction were subjected to agarose gel electrophoresis. The 450 bp band corresponded to the expected length between the two primer sites corresponding to the middle of the third cadherin extracellular subdomain (EC-3) and the carboxyl terminus of the fourth cadherin extracellular subdomain (EC-4), but the 130 bp band could not be predicted from any of the previously identified cadherin sequences. The 450 bp and 130 bp bands were extracted by a freezing and thawing method. The resulting fragments were phosphorylated at the 5' end with T4 polynucleotide kinase and subcloned by a blunt-end ligation into the Sma I site of M13mp18 (Boehringer Mannheim Biochemicals) in a blunt end ligation for sequence analysis. Sequencing of the fragments was carried out by the dideoxynucleotide chain termination method using a Sequenase kit (United States Biochemicals, Cleveland, Ohio). DNA and amino acid sequence were analyzed using the Beckman Microgenie program (Fullerton, California).

Analysis of cDNA Sequences

Nineteen novel partial cDNA clones were isolated. The DNA and deduced amino acid sequences of the clones (including sequences corresponding to the PCR primers) are set out as follows: RAT-123 (SEQ ID NOs: 3 and 4, respectively), RAT-212 (SEQ ID NOs: 5 and 6), RAT-214 (SEQ ID NOs: 7 and

10

15

20

8), RAT-216 (SEQ ID NOs: 9 and 10), RAT-218 (SEQ ID NOs: 11 and 12), RAT-224 (SEQ ID NOs: 13 and 14), RAT-312 (SEQ ID NOs: 15 and 16), RAT-313 (SEQ ID NOs: 17 and 18), RAT-314 (SEQ ID NOs: 19 and 20), RAT-315 (SEQ ID NOs: 21 and 22), RAT-316 (SEQ ID NOs: 23 and 24), RAT-317 (SEQ ID NOs: 25 and 26), RAT-321 (SEQ ID NOs: 27 and 28), RAT-323 (SEQ ID NOs: 29 and 30), RAT-336 (SEQ ID NOs: 31 and 32), RAT-352 (SEQ ID NOs: 33 and 34), RAT-411 (SEQ ID NOs: 35 and 36), RAT-413 (SEQ ID NOs: 37 and 38), and RAT-551 (SEQ ID NOs: 39 and 40).

The deduced amino acid sequences of the cDNA clones are homologous to, but distinct from the known cadherins. The cadherins described thus far have highly conserved, short amino acid sequences in the third extracellular subdomain (EC-3) including the consensus sequence D-Y-E or D-F-E located at the middle region of the subdomain and the consensus sequence D-X-N-E-X-P-X-F (SEQ ID NO: 41) or D-X-D-E-X-P-X-F (SEQ ID NO: 42) at its end (Hatta et al., supra), while the corresponding sequences of other subdomains, except for the fifth extracellular subdomain (EC-5), are D-R-E and D-X-N-D-N-X-P-X-F (SEQ ID NO: 43), respectively. In contrast, the deduced amino acid sequences of the new clones that correspond to cadherin extracellular subdomains include the sequence D-Y-E or D-F-E at one end, but have the instead of D-X-N-E-X-P-X-F sequence D-X-N-D-N-X-P-X-F D-X-D-E-X-P-X-F, at the other end. The polypeptides encoded by the partial clones are homologous to previously identified cadherins but did not show significant homology to any other sequences in Genbank. Therefore, the partial cDNAs appear to comprise a new subclass of cadherin-related molecules.

25

Example 2

Various cDNA fragments structurally similar to the rat cDNAs described in Example 1 were isolated from human, mouse, and Xenopus brain cDNA preparations and from *Drosophila* and *C. elegans* whole body cDNA

10

15

20

25

preparations by PCR using Primers 1 and 2 as described in Example 1. The DNA and deduced amino acid sequences of the resulting PCR fragments (including sequences corresponding to the PCR primers) are set out as follows: MOUSE-321 (SEQ ID NOs: 44 and 45), MOUSE-322 (SEQ ID NOs: 46 and 47), MOUSE-324 (SEQ ID NOs: 48 and 49), MOUSE-326 (SEQ ID NOs: 50 and 51), HUMAN-11 (SEQ ID NOs: 52 and 53), HUMAN-13 (SEQ ID NOs: 54 and 55), HUMAN-21 (SEQ ID NOs: 56 and 57), HUMAN-24 (SEQ ID NOs: 58 and 59), HUMAN-32 (SEQ ID NOs: 60 and 61), HUMAN-42 (SEQ ID NOs: 62 and 63), HUMAN-43 (SEQ ID NOs: 64 and 65), HUMAN-212 (SEQ ID NOs: 66 and 67), HUMAN-213 (SEQ ID NOs: 68 and 69), HUMAN-215 (SEQ ID NOs: 70 and 71), HUMAN-223 (SEQ ID NOs: 72 and 73), HUMAN-410 (SEQ ID NOs: 74 and 75), HUMAN-443 (SEQ ID NOs: 76 and 77), XENOPUS-21 (SEQ ID NOs: 78 and 79), XENOPUS-23 (SEQ ID NOs: 80 and 81), XENOPUS-25 (SEQ ID NOs: 82 and 83), XENOPUS-31 (SEQ ID NOs: 84 and 85), DROSOPHILA-12 (SEO ID NOs: 86 and 87), DROSOPHILA-13 (SEQ ID NOs: 88 and 89), DROSOPHILA-14 (SEQ ID NOs: 90 and 91) and C.ELEGANS-41 (SEQ ID NOs: 92 and 93). Comparison of the deduced amino acid sequences indicates significant similarity between sets of these clones. In particular, there are three sets of clones that appear to be cross-species homologues: RAT-218, MOUSE-322 and HUMAN-43; RAT-314, MOUSE-321 and HUMAN-11; and MOUSE-326 and HUMAN-42.

Example 3

To ascertain the complete structure of the new proteins defined by the PCR products, two full length human cDNAs corresponding to the partial cDNAs HUMAN-42 and HUMAN-43 were isolated.

Isolation of Full-length Human cDNAs

A human fetal brain cDNA library (Stratagene, La Jolla, California) in the λZapII vector was screened by the plaque hybridization method

10

15

20

25

[described in Ausubel et al., Eds., Current Protocols in Molecular Biology, Sections 6.1.1 to 6.1.4 and 6.2.1 to 6.2.3, John Wiley & Sons, New York (1987)] with ³²P-labelled HUMAN-42 and HUMAN-43 DNA fragments. The positive clones were plaque-purified and, using a helper virus, the inserts were cut out by an in vivo excision method in the form of a Bluescript SK(+) plasmid. The insert sequences were then subcloned into the M13 vector (Boehringer Mannheim, Biochemicals) for sequencing. Several overlapping cDNA clones were isolated with each probe including two cDNAs which contained the putative entire coding sequences of two novel proteins designated protocadherin-42 (pc42) and protocadherin-43 (pc43). The DNA and deduced amino acid sequences of pc42 are set out in SEQ ID NOs: 94 and 95, respectively, while the DNA and deduced amino acid sequences of pc43 are set out in SEQ ID NOs: 96 and 97, respectively.

A description of the cloning of protocadherin sequences of the invention was published in Sano et al., The EMBO Journal, 12(6): 2249-2256 (1993) after filing of the priority application hereto. The deduced amino acid sequence of pc43 was previously presented at the December 9, 1991 meeting of the American Society for Cell Biology. An abstract of the presentation is published as Suzuki et al., J. Cell. Biol., 115: 72a (Abstract 416) (December 9, 1991).

Analysis of Full-length Human Clones

Comparison of the full length cDNA sequences of pc42 and pc43 to the sequences of the various DNA fragments originally obtained by PCR reveals that MOUSE-326 and HUMAN-42 correspond to a portion of the fourth extracellular subdomain (EC-4) of pc42, and RAT-314, MOUSE-321, and HUMAN-11 correspond to a portion of the third extracellular subdomain (EC-3) of pc43 and RAT-218, MOUSE-322 and HUMAN-43 correspond to a portion of the fifth extracellular domain (EC-5) of pc43.

10

The overall structures of pc42 and pc43 are similar to that of typical cadherins but the new molecules also have distinct features. Both protocadherin cDNA sequences contain putative translation initiation sites and translated amino acid sequences start with typical signal sequences, but the clones lack the prosequences that are present in all known cadherin precursors. The cDNAs encode proteins having a large N-terminal extracellular domain and a relatively short C-terminal cytoplasmic domain connected by a transmembrane sequence. The extracellular domains of pc42 and pc43 are different in length and pc42 contains seven subdomains that closely resemble the typical cadherin extracellular subdomain while pc43 has six such subdomains. The sizes of the protocadherin cytoplasmic domains are similar to those of typical cadherins, but the sequences do not show any significant homology with those of known cadherins or cadherin-related proteins.

Amino acid identity determinations between extracellular subdomains of human pc42 and pc43, and of mouse N-cadherin (SEQ ID NO: 98) (presented as an example of a "typical" cadherin) and the eighteenth extracellular subdomain of *Drosophila fat* tumor suppressor (EC-18, SEQ ID NO: 99) (the eighteenth extracellular subdomain of *fat* is a prototypical *fat* subdomain) are presented in Table 1 below, wherein, for example, "N-EC-1 x pc42" indicates that the first extracellular subdomain of N-cadherin was compared to the extracellular subdomain of pc42 indicated on the horizonal axis.

20

Table 1

		EC-1	EC-2	EC-3	EC-4	EC-5	EC-6	EC-7
	N-EC-1 x pc42	20	27	26	26	31	29	17
	N-EC-1 x pc43	31	23	23	26	31	24	
5	N-EC-2 x pc42	28	30	32	30	37	31	19
	N-EC-2 x pc43	30	28	30	36	29	30	
	N-EC-3 x pc42	21	26	30	29	31	30	22
	N-EC-3 x pc43	25	18	26	28	28	25	
	N-EC-4 x pc42	28	28	26	25	29	27	17
10	N-EC-4 x pc43	21	25	28	28	29	24	
	N-EC-5 x pc42	24	21	25	24	24	19	12
	N-EC-5 x pc43	15	21	20	20	25	16	
	fat EC-18 x pc42	22	35	32	34	42	35	19
	fat EC-18 x pc43	32	30	36	36	33	29	

The amino acid identity values between the extracellular subdomains of pc42 and pc43, and N-cadherin EC-1 through EC-5 and *Drosophila fat* EC-18 are mostly less than 40%. These identity values are comparable to the values between the subdomains of other cadherin subclasses. However, higher identity values indicate that pc42 and pc43 are more closely related to *fat* than to N-cadherin.

Amino acid identity determinations between extracellular subdomains of human pc42 and pc43 are presented in Table 2 below.

20

Table 2 pc42

<u>pc43</u>	<u>EC-1</u>	<u>EC-2</u>	<u>EC-3</u>	EC-4	<u>EC-5</u>	EC-6	<u>EC-7</u>
EC-1	33	27	29	26	25	26	25
EC-2	26	38	29	33	34	28	21
EC-3	26	32	41	30	32	31	22
EC-4	25	34	30	41	39	31	18
EC-5	23	32	29	27	36	34	16
EC-6	25	25	26	25	28	23	26

The identity values between respective EC-l, EC-2, EC-3, EC-4, EC-5 subdomains and the last subdomains of pc42 and pc43 are generally higher values than values obtained for comparisons of the protocadherins to N-cadherin. These results suggest that pc42 and pc43 are more closely related to one another than they are to classic cadherins.

FIGURE 1A-C presents an alignment of the deduced amino acid sequences of the extracellular subdomains of pc42 (EC-1 through EC-7), pc43 (EC-1 through EC-6), mouse N-cadherin (EC-1 through EC-5) and *Drosophila fat* EC-18. A sequence on a line in FIGURE 1A continues on the same line in FIGURES 1B and 1C. Gaps were introduced to maximize homology. The amino acid residues described by capital letters in the "motif" line are present in more than half of the subdomains of N-cadherin, pc42, pc43 and *Drosophila fat*. The amino acid residues described by small letters in the motif line are less well conserved in human pc42, pc43, and *Drosophila fat*. FIGURE 1A-C shows that many amino acids characteristic of other cadherin extracellular domain repeats are conserved in the pc42 and pc43 sequences, including the cadherin sequence motifs DXD, DRE and DXNDNXPXF (SEQ ID NO: 43), two glycine residues, and one glutamic acid residue. Additionally, pc42 and pc43 share unique features in comparison to N-cadherin. More amino acids at specific sites are conserved

15

10

5

20

10

15

20

25

between pc42 and pc43, such as the DXDXGXN (SEQ ID NO: 100) protocadherin sequence motif near the amino terminus of the pc42 and pc43 subdomains and the AXDXGXP (SEQ ID NO: 101) sequence motif near the carboxyl terminus of the subdomains. Additionally, both protocadherins share regions that do not show significant homology with the typical cadherin motif (of N-cadherin) near the carboxyl terminus of EC-1, in the middle of EC-2 and EC-4, and at the carboxyl terminus of the last repeat. A cysteine residue is located at a similar position in the middle of EC-4 of pc42 and pc43. In general, the extracellular subdomains of pc42 and pc43 are more similar to EC-18 of fat than the extracellular subdomains of N-cadherin.

Possible Alternative Splicing

Sequence analysis of various overlapping protocadherin cDNA clones revealed that some clones contained unique sequences at the 3' end, although the 5' end sequences were identical to other clones. The sequences forming the boundaries of the 3' end regions are consistent with the consensus sequence of mRNA splicing, suggesting that these clones may correspond to alternatively spliced mRNAs. The DNA and deduced amino acid sequences of one possible product of alternative splicing of pc42 mRNA are set out in SEQ ID NOs: 102 and 103. The DNA and deduced amino acid sequences of two possible products of alternative splicing of pc43 mRNA are respectively presented in SEQ ID NO: 104 and 105, and SEQ ID NOs: 106 and 107.

Chromosome Localization

The chromosomal location of the protocadherin 413 gene (SEQ ID NO: 37) and of the pc42 and pc43 genes was determined by conventional methods.

Briefly, C3H/HeJ-gld and Mus spretus (Spain) mice and $[(C3H/HeJ-gld \times Mus \ spretus) \ F_1 \times C3H/HeJ-gld]$ interspecies backcross mice were bred and maintained as previously described in Seldin, et al., J. Exp. Med., 167: 688-693 (1988). Mus spretus was chosen as the second parent in the cross

10

15

20

because of the relative ease of detection of informative restriction fragment length variants (RFLVs) in comparison with crosses using conventional inbred laboratory strains. Gene linkage was determined by segregation analysis.

Genomic DNA isolated from mouse organs by standard techniques was digested with restriction endonucleases and 10μg samples were electrophoresed in 0.9% agarose gels. DNA was transferred to Nytran membranes (Schleicher & Schull, Inc., Keene, NH), hybridized with the appropriate probe at 65°C and washed under stringent conditions, all as previously described in Maniatis et al., supra). To localize the pc42 gene, a mouse sequence probe corresponding to nucleotides 1419 to 1906 of SEO ID NO: 94 was used and for pc43 a rat sequence probe corresponding to nucleotides 1060 to 1811 of SEQ ID NO: 96 was used. To localize the procadherin 413 gene, a probe including the sequence set out in SEQ ID NO: 37 was used. Other clones used as probes in the current study and RFLVs used to detect anonymous DNA loci were all previously described [Chromosome 7, DNA segment, Washington 12 (D7Was12); the parathyroid hormone (Pth); calcitonin (Calc); hemoglobin, β chain (Hbb); metallothionein-I (Mt-I); adenine phosphoribosyltransferase (Aprt); growth hormone receptor (Ghr); prostaglandin E receptor EP2 subtype (Ptgerep2); dihydrofolate reductase-2 (Dhfr2); fibroblast growth factor a (Fgfa); and glucocorticoid receptor-1 (Grl-1)].

Comparison of the haplotype distribution of protocadherin genes with those determined for loci throughout the mouse genome allowed each to be mapped to specific regions of mouse chromosomes. The probability for linkage was >99% and indicated assignment of both the pc42 gene and the pc43 gene was chromosome 18. The assignment of the protocadherin 413 gene was chromosome 7. The region of chromosome 18 to which the pc42 and pc43 genes were mapped corresponds to the ataxia (ax) loci [Burt, Anat. Rec., 196: 61-69 (1980) and Lyon, J. Hered., 46: 77-80 (1955)] and twirler (Tw) loci [Lyon, J. Embryol. Exp. Morphol., 6: 105-116 (1958)], while the region of chromosome

7 to which the protocadherin 413 gene was mapped corresponds to the shaker (sh-1) locus [Kikuchi et al., Acta Oto-Laryngol., 60: 287-303 (1965) and Lord et al., Am. Nat., 63: 453-442 (1929)]. These loci have been implicated as involved in hereditary neural disease in the mouse. This result is consistent with in situ hybridization results (see Example 12) showing that pc42 and pc43 are strongly expressed in the brain and particularly in the cerebellum.

Example 4

Two additional novel human protocadherin cDNAs and one additional novel rat protocadherin cDNA were isolated using rat protocadherin fragments described in Example 1 as probes.

Initially, the rat clone RAT-214 (SEQ ID NO: 7) was used as a probe to screen a rat brain cDNA library (Stratagene, La Jolla, CA). The final washing step was performed twice at 50°C in 0.1X SSC with 0.1% SDS for 15 minutes. Various clones were identified which contained partial cDNA inserts encoding related protocadherin amino acid sequences. The nucleotide sequence of one novel rat clone designated #6-2 is set out in SEQ ID NO: 108. The first fifteen nucleotides of SEQ ID NO: 108 are the sequence of a linker and are not part of the rat #6-2 clone.

A human fetal brain cDNA library obtained from Stratagene was screened with the 0.7 kbp PstI fragment of clone #6-2. The fragment appears to encode the EC-2 and EC-3 of the rat protocadherin. After screening about 2x10⁶ phages, eleven positive clones were isolated. Sequencing of the clones identified a novel full length human protocadherin cDNA designated human pc3. The nucleotide and deduced amino acid sequence of human pc3 are set out in SEQ ID NOs: 109 and 110.

The 0.7 kbp PstI fragment of rat clone #6-2 was also used to rescreen the Stratagene rat brain cDNA library for full length rat cDNA clones. A clone containing an insert encoding a full length novel protocadherin cDNA

15

10

5

20

10

15

20

25

was isolated. The DNA and deduced amino acid sequence of the insert are set out in SEQ ID NO: 111 and 112. The full length rat cDNA was named pc5 because it does not appear to be the homolog of the human pc3 clone based upon a comparison of the sequences.

Concurrently, the 0.8 kbp Eco RI-Pst I fragment of partial rat cDNA designated #43 (SEQ ID NO: 113), which was obtained by screening the Stratagene rat brain cDNA library with a probe corresponding to the human pc43 cytoplasmic domain, was used to probe the Stratagene human cDNA library for full length human protocadherin cDNAs. The fragment appears to encode EC-3 through the beginning of EC-6 of clone #43. One partial clone identified encodes a novel human protocadherin named human pc4. The nucleotide sequence and deduced amino acid sequences of the human pc4 clone are set out in SEQ ID NOs: 114 and 115. The amino acid sequence encoded by the pc4 clone appears to begin in the middle of EC-2 of pc4 and continues through the cytoplasmic tail of the protocadherin.

Example 5

The full length human cDNAs encoding pc42 and pc43 were expressed in L cells (ATCC CCL 1) using the pRC/RSV expression vector (Invitrogen, San Diego, California). The cDNAs were isolated from the Bluescript SK(+) clones described in Example 2 by digestion with SspI followed by blunt-ending with DNA polymerase and digestion with XbaI (for pc42), or by double digestion with SpeI and EcoRV (for pc43). The pRC/RSV expression vector was digested with HindIII, followed by blunt-ending and re-digestion with XbaI for insertion of pc42 sequences, or by digested with XbaI followed by blunt-ending and re-digestion with SpeI for insertion of pc43 sequences. The isolated protocadherin DNAs were ligated into the linearized pRC/RSV vector. The resulting pc42 expression plasmid designated pRC/RSV-pc42 (ATCC 69162) and pc43 expression plasmid designated pRC/RSV-pc43 (ATCC 69163) were

10

15

20

purified by CsCl gradient centrifugation and transfected into L cells by a Caphosphate method.

The pc42 and pc43 transfectants were morphologically similar to the parental cells. Northern blot analysis of L cells transfected with pc42 or pc43 DNA sequences showed that the transfected cells expressed mRNAs of a size expected to encode the particular protocadherin.

Example 6

Rabbit polyclonal antibodies specific for pc42 and pc43 were generated as well as a mouse monoclonal antibody specific for pc43.

Preparation of Polyclonal Antibodies Specific for pc42 and pc43

DNA sequences encoding portions of the extracellular domain of pc42 and pc43 were each fused to a maltose binding protein-encoding sequence and expressed in bacteria. Specifically, DNAs corresponding to EC-4 through EC-7 of pc42 and EC-3 through EC-5 of pc43 were prepared by PCR and subcloned in the correct reading frame into the multicloning site of the pMAL expression vector (New England Biolabs, Beverly, Massachusetts) which contains sequences encoding maltose binding protein immediately upstream of the multicloning site. The resulting plasmids were then introduced into E. coli NM522 cells (Invitrogen, San Diego, California) by a single step transformation method. Expression of the fusion proteins was induced by the addition of IPTG and the fusion proteins were purified from cell extracts by amylose resin affinity chromatography (New England Biolabs) as described by the manufacturer. The fusion proteins were used for the immunization of rabbits without further purification.

25

Polyclonal antibodies were prepared in rabbits by immunization at four subcutaneous sites with $500\mu g$ of purified fusion protein in Freund's complete adjuvant. Subsequent immunizations with $100\mu g$ of the fusion protein were in Freund's incomplete adjuvant. Immune sera was passed through

10 -

15

sepharose coupled to maltose binding protein (New England Biolabs) and polyclonal antibodies were purified from immune sera using Sepharose affinity columns prepared by reaction of the purified fusion protein with CNBr Sepharose (Pharmacia). Reactivity of the polyclonal sera with purified pc42 fusion protein and pc42 transfected cell extracts (described in Example 5) was confirmed.

Preparation of Monoclonal Antibodies Specific for pc43

The pc43 fusion protein (containing the EC-3 through EC-5 subdomains of pc43) was used to generate monoclonal antibodies in mice according to the method of Kennett, *Methods in Enzymol.*, 58:345-359 (1978). Briefly, mice were immunized with the pc43 fusion protein ($100\mu g$) at two subcutaneous sites. The spleen from the highest titer mouse was fused to the NS1 myeloma cell line. The resulting hybridoma supernatants were screened in a ELISA assay for reactivity with the pc43 fusion protein and with maltose binding protein. The fusion wells with the highest reactivity to the pc43 extracellular domains were subcloned. The hybridoma cell line designated 38I2C (ATCC HB 11207) produced a IgG_1 subtype monoclonal antibody specific for pc43. Reactivity of the monoclonal antibody produced by hybridoma cell line 38I2C to pc43 was confirmed by immunoblotting the pc43 L cell transfectants described in Example 5. The 38I2C monoclonal antibody is specific for human pc43.

20

Example 7

L cells transfected with DNA sequences encoding pc42 and pc43 as prepared in Example 5 were assayed for expression of the protocadherins by immunoblot and by immunofluorescence microscopy.

Immunoblot Analysis

25

Cell extracts of pc42 and pc43 transfectants were subjected to SDS-PAGE and then blotted electrophoretically onto a PVDF membrane (Millipore, Bedford, Massachusetts). The membranes were incubated with 5% skim milk in Tris-buffered saline (TBS) for two hours and then respectively with

10

15

20

25

either pc42 polyclonal sera or pc43 monoclonal antibody for one hour. The membranes were washed three times (for 5 minutes each wash) with TBS containing 0.05% Tween 20 and respectively incubated with alkaline phosphatase-conjugated anti-rabbit IgG antibody or anti-mouse IgG antibody (Promega, Madison, Wisconsin) in the same buffer for one hour. After washing the membranes with TBS containing 0.05% Tween 20, reactive bands were visualized by using Western Blue solution (Promega).

Anti-pc42 polyclonal antibodies stained a band of about 170 kDa molecular weight in pc42 transfected cells, but not parental L cells. The pc43-specific monoclonal antibody (38I2C) and polyclonal antibodies stained two adjacent bands of about 150 kDa molecular weight in pc43 transfected cells. The pc43 antibodies did not stain bands in parental L-cells. The molecular weights indicated by the staining of bands by the pc42 and pc43 antibodies are significantly larger than the molecular weights predicted from the deduced amino acid sequences. This discrepancy in molecular weight is common among various cadherin-related proteins and may be attributable to the glycosylation and/or cadherin specific structural properties. The pc42 antibody also stained smaller bands, which may be proteolytic degradation products.

When transfected cells were trypsinized and cell extracts were prepared, run on SDS/PAGE and immunoblotted with the appropriate antibody, the pc42 and pc43 polypeptides expressed by the transfected cells were found to be highly sensitive to proteolysis and were easily digested by 0.01% trypsin treatment. In contrast to the classic cadherins, however, these proteins were not protected from the digestion in the presence of 1-5mM Ca²⁺.

Immunofluorescence Microscopy

Transfected cells were grown on a cover slip precoated with fibronectin and were fixed with 4% paraformaldehyde for 5 minutes at room temperature or with cold methanol on ice for 10 minutes followed by 4% paraformaldehyde fixation. After washing with TBS, the cells were incubated with

10

15

20

TBS containing 1% BSA for 30 minutes and then with anti-pc42 polyclonal antibody or anti-pc43 monoclonal antibody in TBS containing 1% BSA for 1 hour at room temperature. Cover slips were then washed with TBS containing 0.01% BSA and respectively incubated with FITC-conjugated anti-rabbit antibody or anti-mouse antibody (Cappel, Durham, North Carolina) for 60 minutes at room temperature. The cells were washed again with TBS containing 0.01% BSA and subjected to fluorescence microscopy. Both pc42-specific and pc43-specific polyclonal antibodies stained the cell periphery of transfected cells expressing the protocadherin proteins, mainly at the cell-cell contact sites. The antibodies did not stain the parent L cells, nor did rabbit preimmune sera stain the pc42 and pc43 transfectants.

Example 8

The cell aggregation properties of the transfected L cells expressing protocadherin proteins were examined. Transfected L cells were cultured in Dulbecco's Modified Eagles Medium (DMEM) (Gibco, Grand Island, New York) supplemented with 10% fetal bovine serum at 37°C in 5% CO₂. Cells grown near confluence were treated with 0.01% trypsin in the presence of 1 mM EGTA for 25 minutes on a rotary shaker at 37°C and collected by centrifugation. The cells were washed three times with Ca²⁺ free HEPES-buffered saline (HBS) after adding soybean trypsin inhibitor, and were resuspended in HBS containing 1% BSA. The cell aggregation assay [Urushihara et al., Dev. Biol., 70: 206-216 (1979)] was performed by incubating the resuspended cells in a 1:1 mixture of DMEM and HBS containing 1% BSA, 2 mM CaCl₂ and 20 µg/ml of deoxyribonucelease on a rotary shaker at 37°C for 30 minutes to 6 hours.

The pc42 and pc43 transfectants did not show any significant cell aggregation activity during periods of incubation less than 1 hour. This is in contrast to the cell aggregation that occurs with classic cadherins in similar experiments (Nagafuchi et al., supra, and Hatta et al., supra). However,

10

15

20

25

30

prolonged incubation of transfected cells (more than 1-2 hours) resulted in gradual re-aggregation of the cells into small aggregates. Similar results were obtained when single cell suspensions of transfected cells were prepared by trypsin treatment in the presence of Ca²⁺. No re-aggregation was observed under the same conditions when untransfected L cells or L cells transfected with pRC/RSV vector alone were tested. When pc43 transfectants labelled with DiO (Molecular Probes, Eugene, OR) were incubated with unlabelled pc42 transfectants in the cell aggregation assay, aggregation of labelled and unlabelled cells was almost mutually exclusive indicating that protocadherin binding is homophilic.

In view of the fact that the protocadherin cytoplasmic domains exhibit no apparent homology to cadherin domains, experiments were performed to determine if the difference in cytoplasmic domains could account for the difference in cell aggregation activity observed in cadherin and protocadherin transfectants. The cytoplasmic domain of pc43 was replaced with the cytoplasmic domain of E-cadherin and aggregation of cells transfected with the chimeric construct was analyzed.

The Bluescript SK(+) clone described in Example 2 which contained the entire coding sequence for pc43 was digested with EcoRV and then partially digested with XbaI to remove the sequence corresponding to the cytoplasmic domain, and the plasmid DNA was purified by agarose gel electrophoresis. The cDNA corresponding to the cytoplasmic domain of mouse E-cadherin was synthesized by PCR using mouse cDNA made from mouse lung mRNA as a template and specific primers corresponding to a region near the N-terminus of the cytoplasmic domain sequence or the region containing the stop codon of mouse E-cadherin (Nagafuchi et al., supra). A XbaI sequence was included to the 5' end of the upstream primer. The E-cadherin cytoplasmic domain cDNA was then subcloned into the linearized pc43 Bluescript clone. The DNA containing the entire resulting chimeric sequence was cut out with SpeI and EcoRV and was subcloned into the SpeI-blunted XbaI site of the expression vector pRc/RSV vector. Finally, L cells were transfected with the resultant construct by

10

15

25

a calcium phosphate method. After screening with G418 for about 10 days, the transfectants were stained with FITC-labeled 38I2C anti-pc43 antibody and subjected to FACS analysis. A portion of highly labeled cells were isolated and cloned. Transfectants showed a morphology similar to that of parental L cells and the expressed protein was localized at the cell periphery using pc43 antibody for immunofluorescence microscopy.

Cell aggregation activity of the chimeric transfectants was analyzed as follows. The chimeric pc43 transfectants were labeled with DiO for 20 minutes at room temperature. The resultant cells were trypsinized in the presence of 1mM EGTA and single cell suspension was made. Then, the cells were mixed with unlabeled other type of transfectants and incubated on a rotary shaker for two hours. The results were examined with a fluorescence and a phase contrast microscope apparatus. Antibody inhibition of cell aggregation was examined by incubation of the transfectants in the presence of polyclonal anti-pc43 antibody (100 ng/ml) in the standard assay medium.

In the cell aggregation assay, the chimeric pc43 transfectants showed clear Ca²⁺-dependent cell aggregation within forty minutes of incubation. Cell aggregation was inhibited by the addition of pc43-specific polyclonal antibody.

20 Example 9

The procedures of Maruyama et al., J. Biochem., 95: 511-519 (1984) were used to determine the calcium binding properties of pc43 by Western blot analysis in the presence or absence of calcium-45. The pc43 fusion protein described in Example 6 containing pc43 subdomains EC-3 through EC-5 was compared to the calcium binding protein calmodulin. Samples of purified pc43 fusion protein were run on SDS/PAGE and electrophoretically transferred to PVDF membrane. Binding of the ⁴⁵Ca²⁺ to the pc43 fusion protein was detected by autoradiography and was determined to be nearly as efficient as binding of ⁴⁵Ca²⁺ to calmodulin. In contrast, there was no binding of calcium to purified

maltose binding protein lacking the pc43 extracellular domain. The pc43 subdomains EC-3 through EC-5 contain sequences highly homologous to the putative $Ca^{2\pm}$ binding motifs found in E-cadherin. [See, Ringwald *et al.*, *EMBO J.*, 6: 3647-3653 (1987).]

5

10

15

20

25

Example 10

The expression of mRNA encoding pc42 and pc43 was assayed in various tissues and cell lines by Northern blot.

Total RNAs were prepared by the guanidium isothiocyanate method and poly(A)+ RNAs were isolated using a FastTrack kit (Invitrogen). RNA preparations were electrophoresed in a 0.8% agarose gel under denaturing conditions and transferred onto a nitrocellulose filter using a capillary method. Northern blot analyses were performed according to the method of Thomas, *Proc. Natl. Acad. Sci. USA*, 77: 5201-5205 (1980). The final wash was in 0.2X standard saline citrate containing 0.1% sodium dodecyl sulfate at 65°C for 10 minutes.

Protocadherin mRNA Expression in Adult Rat Tissues

Total mRNA preparations of rat tissues including brain, heart, liver, lung, skin, kidney and muscle were separated electrophoretically under denaturing conditions (10 μ g mRNA/lane) and transferred onto nitrocellulose filters. The filters were hybridized with ³²P-labelled cDNA fragments MOUSE-326 (which corresponds to EC-4 of human pc42) and RAT-218 (which corresponds to EC-5 of human pc43). The mRNAs of both protocadherins were highly expressed in brain. The pc42 probe detected a major band of 7 kb and a minor band of 4 kb in size, possibly representing the products of alternative splicing. The pc43 probe hybridized to a major band of 5 kb in size and with minor bands of smaller sizes.

Developmental Expression of Protocadherin mRNA in Rat Brain

To examine the developmental regulation of mRNA expression of the protocadherins, brain mRNA from rats at embryonic days 17 and 20, neonatal

10

15

20

25

days 5 and 11 and from adult rats was prepared and subjected to Northern blot analysis as described above for other rat tissues. β -actin was used as an internal standard. mRNA levels for pc42 and pc43 proteins increased during embryonic development of the brain as compared with β -actin expression.

Protocadherin mRNA Expression in Human Cell Lines

Several neuronal and glial cell lines (including human SK-N-SH neuroblastoma, human U251 glioma, and mouse Neuro-2a neuroblastoma cell lines) were assayed by Northern blot using ³²P-labelled for expression of pc42 and pc43 mRNA. Human cell lines were probed with HUMAN-42 (which corresponds to EC-4 of human pc42) and HUMAN-43 (which corresponds to EC-5 of human pc43) cDNA fragments while the mouse cell line was probed with MOUSE-326 (which corresponds to EC-4 of human pc42) and RAT-322 (which corresponds to EC-5 of human pc43) cDNA fragments. SK-N-SH human neuroblastoma cells and U251 human glioma cells were found to express pc43 mRNA and Neuro-2a mouse neuroblastoma cells were found to express pc42 mRNA.

Example 11

Expression of pc43 protein in various tissues, extracts and cells was assayed by Western blot and immunofluorescence microscopy.

Expression in Rat Cardiac Muscle Extracts

A rat heart non-ionic detergent extract was prepared by freezing a heart in liquid nitrogen after removal, powdering in a mortar and pestle, grinding briefly in a polytron in 0.5% Nonidet P40 in [10 mM PIPES (pH 6.8), 50 mM NaCl, 250 mM NH₄SO₄, 300 mM sucrose, 3 mM MgCl₂] and microfuging for 15 minutes. Samples were separated by SDS/PAGE and electrophoretically transferred to nitrocellulose (Towbin *et al.*, *PNAS 76:*4350-4354, 1979). Two pc43 protein bands with molecular weights of 150 KDa and 140 KDa were

10

15

detected with rabbit polyclonal antibodies to pc43 by the immunoblot method described in Example 7.

Expression in Tissue Sections and Cells

To determine the localization of the protocadherins in various tissues, human and rat adult tissues were removed, incubated in 30% sucrose in PBS for 30 minutes at 4°C, embedded in OCT compound (Tissue-Tek, Elkhart, Indiana) in cryomolds and quickly frozen. Six micron sections were cut and placed on glass slides. The slides were washed with PBS and fixed in 3% p-formaldehyde for 5 minutes. To permeablize the tissue sections, the slides were immersed in -20°C acetone for 10 minutes and air dried. The sections were blocked with 2% goat serum and 1% BSA in PBS for 30 minutes and then incubated with the rabbit anti-pc43 polyclonal antisera for 1 hour at room temperature. The sections were rinsed 3 times in PBS containing 0.1% BSA and incubated with a biotinylated anti-rabbit (Vector Laboratories, Burlingame, California) in 1% BSA in PBS for 30 minutes. After rinsing 3 times, strepavidinconjugated with FITC (Vector Laboratories) was added for 30 minutes and again washed 3 times. For co-localization studies, an appropriate primary antibody was used with a TRITC-conjugated secondary antibody.

A. Muscle

20

25

Immunolocalization of pc43 in rat cardiac muscle shows that pc43 is localized in a repeating pattern which is consistent with pc43 being associated with the sarcomeres. Sarcomeres are repetitive contractile units between the fascia adherens in skeletal and cardiac muscle. Co-localization with cytoskeletal proteins shows that pc43 is present at the ends of the sarcomeres in the Z lines which are associated with desmin and the actin-binding protein vinculin, and alpha-actinin. The thin microfilaments of F-actin are associated with the thick myosin filaments between the Z lines. In contrast, N-cadherin is localized at the ends of cardiac myocytes at the fascia adherens junctions at sites of mycocyte:myocyte contact. The localization of pc43 in cardiac muscle suggests

10

15

20

25

that pc43 may play a role in muscle contraction in the anchoring of the contractile apparatus to the plasma membrane.

Similar localization for pc43 was observed in rat skeletal muscle. Ultrastructural studies have shown that dystrophin, the gene product lacking in Duchenne muscular dystrophy, is a component of the sarcolemma [Porter et al., J. Cell. Biol., 117:997-1005 (1992)]. The sarcolemma is connected to the contractile apparatus at the M and Z lines where pc43 is localized.

B. Brain

Reactivity of anti-pc43 polyclonal antibody and monoclonal antibody 38I2C on frozen sections of rat and human cerebellum, respectively, shows that the major sites of pc43 expression are located in Purkinje cells and the granule cell layer which contains numerous small neurons.

C. Placenta

Strong reactivity of monoclonal antibody 38I2C with human syncytiotrophoblasts was also observed in development of the placenta at an early state (5-7 weeks of gestation). Expression appeared to gradually decrease as the stage progressed indicating that pc43 may be involved in the implantation of fertilized eggs into the placenta.

D. Neuroblastoma and Astrocytoma Cells

Immunocytochemical localization of pc43 in Sk-N-SH neuroblastoma cells and UW28 astrocytoma cells using anti-pc43 antibodies reveals a punctate cell surface distribution of pc43 and in some cells there is a localization at the tips of extensions of neuronal foot processes. At sites of cell-cell contact of UW28 astrocytoma cells, pc43 is organized in a series of parallel lines. The lines start at the contact site and extend approximately 5 micron. Factin microfilaments were identified with rhodamine-phalloidin (Molecular Probes, Eugene, Oregon, as described by the manufacturer) showing that the microfilaments in the cell appear to end in the pc43 linear structures which extend from the edge of the cell at sites of cell contact.

Immunoblotting studies with pc43 specific antibodies show that a protein with a molecular weight of 140 kDa is recognized in human Sk-N-SH neuroblastoma cells and in UW28 astrocytoma cells.

E. Osteoblasts

5

Immunocytochemical localization of pc43 using monoclonal antibody 38I2C in tow human ostogenic sarcoma cell lines [SaOS (ATCC HTB 85) and MG-63 (ATCC CRL 1427)] and in cultures of normal human trabecular osteoblasts [culture system described in Civitelli et al., J. Clin. Invest., 91: 1888-1896 (1993)] showed that pc43 is expressed in osteoblasts in a pattern similar to that seen in UW28 astrocytoma cells. At sites of cell-cell contact, pc43 is organized in a series of parallel lines that appear to correspond to the actin stress fibers. In addition, in some cells, pc43 appears to localize at the tips of contacting cell processes. Northern blot analysis provides additional evidence that pc43 is expressed in normal human trabecular osteoblasts. A pc43 specific DNA probe hybridized to a major band of 5 kb in samples of poly-A mRNA isolated from normal human trabecular osteoblasts.

15

10

Example 12

In situ hybridization experiments using protocadherin specific RNA probes were performed on cryosections of rat tissue.

20

Sense and antisense ³⁵S-riboprobes were made using the standard procedure described by Promega (Madison, Wisconsin). An approximately 400 bp EcoRI-Xbal fragment of the MOUSE-326 cDNA clone was used as a pc42 specific probe. This fragment encodes the middle of EC-3 to the end of EC-4 of pc42. An approximately 700 bp Smal fragment of the RAT-218 cDNA clone was used as a pc43 specific probe. The fragment encodes the end of EC-3 to the end of EC-5 of pc43.

25

Rat adult tissues were harvested and immediately embedded with OCT Compound (Tissue-Tek) in cryomolds and quickly frozen in a bath of 95% ethanol/dry ice. The frozen blocks were stored at -80°C until cut. Six micron

10

15

20

25

tissue sections were cut using a cryostat (Reichert-Jung, Model #2800 Frigocut N, Leica, Inc., Gilroy, California). Cut tissue sections were stored at -80°C.

The in situ protocol used was a variation of that described by Angerer et al., Methods in Enzymology, 152: 649-660, (1987). All solutions were treated with diethylpyrocarbonate (DEPC, Sigma, St. Louis, Missouri) to remove RNase contamination. The tissue sections were first fixed in 4% paraformaldehyde at 4°C for 20 minutes. To remove excess paraformaldehyde and stop the tissue fixation, the slides were washed in PBS (phosphate buffered saline), denatured in a graded series of alcohols (70, 95, 100%) and then dried. To prevent the tissue from detaching from the glass slide during the in situ procedure, the tissue sections were treated in a poly-L-lysine solution (Sigma) at room temperature for 10 minutes. To denature all RNA in the tissue, the sections were placed in a solution of 70% formamide/2x SSC (0.15 M NaCl/0.3 M Na citrate, pH 7.0) at 70°C for 2 minutes after which they were rinsed in chilled 2x SSC, dehydrated in a graded series of alcohols and then dried. Once dried, the sections were prehybridized in hybridization buffer [50% formamide/50 mM DTT (dithiothrietol)/0.3M NaCl/20 mM Tris, pH 8.0/5 mM EDTA/1X Denhardt's (0.02% Ficoll Type 400/0.02% polyvinylpyrrolidone/0.02% BSA)/10% Dextran Sulfate] at the final hybridization temperature for approximately 4 hours. After prehybridization, approximately 1 X 106 cpm of the appropriate riboprobe was added to each section. The sections were generally hybridized at 45°C overnight (12-16 hours). To insure that the hybridization seen was specific, in some experiments the hybridization stringency was increased by raising the hybridization temperature to 50°C. As both the 45°C and 50°C experiments gave comparable results, the standard hybridization temperature used was 45°C.

To remove excess, nonhybridized probe, the sections were put through a series of washes. The sections were first rinsed in 4X SSC to remove the bulk of the hybridization solution and probe. Next a 15 minute wash in 4X SSC/50 mM DTT was carried out at room temperature. Washes at increased

stringencies were also utilized. A 40 minute wash in 50% formamide/2X SSC/50 mM DTT was performed at 60°C. Four final room temperature washes were carried out for 10 minutes each: two in 2X SSC and two in 0.1X SSC. The washed slides were dehydrated in a graded series of alcohols and dried.

5

To visualize the hybridized probe, the slides were dipped in Kodak NTB2 nuclear emulsion (International Biotechnology, New Haven, Connecticut) which had been diluted 1:1 in dH₂O. Once dry, the slides were stored at 4°C in light-tight boxes for the appropriate exposure time. The *in situ* slides were independently viewed by two persons and scored positive or negative for hybridization signal.

10

15

20

All in situ hybridization studies were performed on rat tissue. Because results from Northern blot experiments (see Example 9) indicated that both pc42 and pc43 are expressed in adult brain, in situ hybridization studies were carried out to localize the expression of these molecules to specific brain cell types. Hybridization seen in the normal adult rat brian was specific (no background hybridization was seen with the sense probes) and was localized to specific regions in the brain. The overall pattern of expression seen for pc42 and pc43 was very similar, with the major difference being in the level of expression. pc43 appears to be expressed at a lower level than pc42. Both molecules are expressed in the germinal and pyramidal cells of the hippocampus, Purkinje cells of the cerebellum and neurons in grey matter. In addition, pc42 is expressed in glial cells in the white matter but, in contrast to the expression of pc43 in glioma cell lines (as described in Example 9), expression of pc43 in normal glial cells was not observed. In the spinal chord, both protocadherins are expressed in the motor neurons in the gray matter and pc42 is expressed in the glial cells in the white matter.

25

When expression of both protocadherin molecules was analyzed in brains and spinal chords from rats having EAE (experimental allergic encephalomyelitis) [Vandenbark et al., Cell. Immunol., 12: 85-93 (1974)], the same structures as described above were found to be positive. In addition,

10

15

20

25

expression of pc42 was observed in the leukocytic infiltrates in the EAE tissues. Expression of pc42 in leukocytes was confirmed by *in situ* hybridization analysis of two leukocytic cell lines, RBL-1 and y3.

Expression of both protocadherin-42 and -43 was observed in the developing brain of rat embryos at all embryological days tested (E15-E19). In addition protocadherin-43 was observed in the developing rat heart at all embryological days tested (E13-E19). This finding is consistent with the immunohistochemistry results showing protocadherin-43 expression in adult heart.

To determine possible roles of protocadherins in the development of the nervous system, expression profiles of protocadherin members in developing rat brain and adult rat brain were also examined by in situ hybridization. A series of coronal, sagittal and horizontal sections of rat brains at postnatal days 0, 6, 14, 30 (P0 through P30) and at 3 months (young adult) were hybridized with labelled cRNA probes corresponding to various protocadherins of the invention including pc42, pc43, RAT-212, RAT-411, and RAT-418. In developing brain, RAT-411 was expressed at high levels in neurons of the olfactory bulb, i.e., mitral cells and periglomerular cells. The expression of RAT-411 mRNA was transient; expression appeared at P0, peaked at P6, diminished by P14, and was undetectable at P30 and in adult brain. In the adult, pc43 mRNA was found to be expressed predominantly in Purkinje cells in the cerebellum. The expression of pc43 mRNA in Purkinje cells was observed from the beginning of Purkinje cell differentiation at around P6. Other protocadherin members were expressed at very low levels in various areas of developing and adult brains. These results indicate that protocadherin members are differentially expressed during the development of the central nervous system, and suggest that RAT-411 and pc43 have specific roles during the development of olfactory bulb neurons and Purkinje cells, respectively.

Example 13

Conventional immunoprecipitations using pc43-specific polyclonal antibodies and monoclonal antibody 38I2C were performed to identify proteins that interacted with pc43 in L cell transfectants.

5

The pc43 and chimeric pc43 transfectants were metabolically labeled by incubating the cells in Dulbecco's modified Eagle's medium containing [35S] methionine (50 uCi/ml) overnight. After washing, the transfectants were lysed with PBS containing Triton X 100 and incubated with anti-pc43 antibody. The immunocomplexes were then collected using protein A-Sepharose beads. The resulting beads were washed five times with a washing buffer (50mM Tris-HCl, pH 8.0, containing 0.5M NaCl, 0.1% ovalbumin, 0.5% NP-40, 0.5% Triton X 100 and 1mM EDTA) at room temperature. Protein was separated by SDS-PAGE and subjected to autoradiography.

10

The chimeric pc43 co-precipitated with 105 kDa and a 95 kDa bands that are likely to correspond to α - and β -catenins, respectively, because anti- α -catenin and anti- β -catenin antibodies stained comparable bands. Pc43, on the other hand, co-precipitated with a 120 kDa band.

15

While the present invention has been described in terms of specific methods and compositions, it is understood that variations and modifications will occur to those skilled in the art. Therefore, only such limitations as appear in the claims should be placed on the invention.

SEQUENCE LISTING

- (1) GENERAL INFORMATION:
 - (i) APPLICANT: Suzuki, Shintaro
 - (ii) TITLE OF INVENTION: Protocadherin Materials and Methods
 - (iii) NUMBER OF SEQUENCES: 115
 - (iv) CORRESPONDENCE ADDRESS:
 - (A) ADDRESSEE: Marshall, O'Toole, Gerstein, Murray, & Borun
 - (B) STREET: 6300 Sears Tower, 233 S. Wacker Drive
 - (C) CITY: Chicago
 - (D) STATE: Illinois
 - (E) COUNTRY: USA
 - (F) ZIP: 60606
 - (v) COMPUTER READABLE FORM:
 - (A) MEDIUM TYPE: Floppy disk
 - (B) COMPUTER: IBM PC compatible
 - (C) OPERATING SYSTEM: PC-DOS/MS-DOS
 - (D) SOFTWARE: PatentIn Release #1.0, Version #1.25
 - (vi) CURRENT APPLICATION DATA:
 - (A) APPLICATION NUMBER:
 - (B) FILING DATE:
 - (C) CLASSIFICATION:
 - (vii) PRIOR APPLICATION DATA
 - (A) APPLICATION NUMBER: PCT/US93/12588
 (B) FILING DATE: 23 DEC 1993
 - (vii) PRIOR APPLICATION DATA
 - (A) APPLICATION NUMBER: US 07/998,003
 - (B) FILING DATE: 29 DEC 1992
 - (viii) ATTORNEY/AGENT INFORMATION:

 - (A) NAME: Noland, Greta E. (B) REGISTRATION NUMBER: 35,302
 - (C) REFERENCE/DOCKET NUMBER: 32149
 - (ix) TELECOMMUNICATION INFORMATION:
 - (A) TELEPHONE: 312/474-6300
 - (B) TELEFAX: 312/474-0448
 - (C) TELEX: 25-3856
- (2) INFORMATION FOR SEQ ID NO:1:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 17 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

AARSSNNING AYTRYGA	17
(2) INFORMATION FOR SEQ ID NO:2:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 17 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: DNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:	
TTRCTRTTRC GNGGNNN	17
(2) INFORMATION FOR SEQ ID NO:3:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:	
AAGGGAGTGG ACTTTGAGGA GCAGCCTGAG CTTAGTCTCA TCCTCACGGC TTTGGATGGA	60
GGGACTCCAT CCAGGTCTGG GACTGCATTG GTTCAAGTGG AAGTCATAGA TGCCAATGAC	120
AACGCACCGT A	131
(2) INFORMATION FOR SEQ ID NO:4:	
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 	
(ii) MOLECULE TYPE: protein	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:	
Lys Gly Val Asp Phe Glu Glu Gln Pro Glu Leu Ser Leu Ile Leu Thr 1 5 10 15	
Ala Leu Asp Gly Gly Thr Pro Ser Arg Ser Gly Thr Ala Leu Val Gln 20 25 30	
Val Glu Val Ile Asp Ala Asn Asp Asn Ala Pro 35 40	

(2) INFORMATION FOR SEQ ID NO:5:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:	
AAACGCATGG ATTTCGAGGA GTCTTCCTCC TACCAGATCT ATGTGCAAGC TACTGACCGG	6
GGACCAGTAC CCATGGCGGG TCATTGCAAG GTGTTGGTGG ACATTATAGA TGTGAACGAC	12
AACGCACCTA A	13
(2) INFORMATION FOR SEQ ID NO:6:	
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 	
(ii) MOLECULE TYPE: protein	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:	
Lys Ala Met Asp Phe Glu Glu Ser Ser Ser Tyr Gln Ile Tyr Val Gln 1 5 10 15	
Ala Thr Asp Arg Gly Pro Val Pro Met Ala Gly His Cys Lys Val Leu 20 25 30	
Val Asp Ile Ile Asp Val Asn Asp Asn Ala Pro 35 40	
(2) INFORMATION FOR SEQ ID NO:7:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:	

AAGCGACTGG ACTTTGAGAC CCTGCAGACC TTCGAGTTCA GCGTGGGTGC CACAGACCAT

GGCTCCCCCT CGCTCCGCAG TCAGGCTCTG GTGCGCGTGG TGGTGCTGGA CCACAATGAC

60

120

AATGCCCC	CA A	131
(2) INFO	RMATION FOR SEQ ID NO:8:	
(i)	SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii)	MOLECULE TYPE: protein	
(×i)	SEQUENCE DESCRIPTION: SEQ ID NO:8:	
Lys 1	Arg Leu Asp Phe Glu Thr Leu Gln Thr Phe Glu Phe Ser Val Gly 5 10 15	
Ala	Thr Asp His Gly Ser Pro Ser Leu Arg Ser Gln Ala Leu Val Arg 20 25 30	
Val	Val Val Leu Asp His Asn Asp Asn Ala Pro 35 40	
(2) INFO	RMATION FOR SEQ ID NO:9:	
(i)	SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii)	MOLECULE TYPE: cDNA	
(×i)	SEQUENCE DESCRIPTION: SEQ ID NO:9:	
AAGGGCCT	GG ATTACGAGGC ACTGCAGTCC TTCGAGTTCT ACGTGGGCGC TACAGATGGA	60
GGCTCACC	CG CGCTCAGCAG CCAGACTCTG GTGCGGATGG TGGTGCTGGA TGACAACGAC	120
AACGCCCC	TA A	131
(2) INFO	RMATION FOR SEQ ID NO:10:	
(i)	SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii)	MOLECULE TYPE: protein	

Lys Gly Leu Asp Tyr Glu Ala Leu Gln Ser Phe Glu Phe Tyr Val Gly 1 5 15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:-

Ala	Thr	Asp	Gly	Gly	Ser	Pro	Ala	Leu	Ser	Ser	Gln	Thr	Leu	Val	Arc
			20					25					30		_

Met Val Val Leu Asp Asp Asn Asp Asn Ala Pro 35

- (2) INFORMATION FOR SEQ ID NO:11:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 131 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

AAGGCGTTTG ATTTTGAGGA TCAGAGAGAG TTCCAGCTAA CCGCTCATAT AAACGACGGA 60
GGTACCCCGG TTTTGGCCAC CAACATCAGC GTGAACATAT TTGTTACTGA CCGCAATGAC 120
AACGCCCCGC A 131

- (2) INFORMATION FOR SEQ ID NO:12:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 43 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

Lys Ala Phe Asp Phe Glu Asp Gln Arg Glu Phe Gln Leu Thr Ala His

Ile Asn Asp Gly Gly Thr Pro Val Leu Ala Thr Asn Ile Ser Val Asn 20 25 30

Ile Phe Val Thr Asp Arg Asn Asp Asn Ala Pro 35 40

- (2) INFORMATION FOR SEQ ID NO:13:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 131 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:											
AAGGCGGTGG ATTACGAAAT CACCAAGTCC TATGAGATAG ATGTTCAAGC CCAAGATCTG	60										
GGTCCCAATT CTATTCCTGC TCATTGCAAA ATTATAATTA AGGTCGTGGA TGTCAACGAC	120										
AACGCTCCCA A	131										
(2) INFORMATION FOR SEQ ID NO:14:											
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear (ii) MOLECULE TYPE: protein											
(a=, assessed 22221 p20021											
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:											
Lys Ala Val Asp Tyr Glu Ile Thr Lys Ser Tyr Glu Ile Asp Val Gln											
1 5 10 15											
Ala Gln Asp Leu Gly Pro Asn Ser Ile Pro Ala His Cys Lys Ile Ile 20 25 30											
Ile Lys Val Val Asp Val Asn Asp Asn Ala Pro 35 40											
(2) INFORMATION FOR SEQ ID NO:15:											
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 135 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear											
(ii) MOLECULE TYPE: cDNA											
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:											
TATGACCATG ATTACGAGAC AACCAAAGAA TATACACTGC GGATCCGGGC CCAGGATGGT	60										
GGCCGGACTC CACTTTCCAA CGTCTCCGGT CTAGTAACCG TGCAGGTCCT AGACATCAAC	120										
GACAATGCCC CCCCA	135										
(2) INFORMATION FOR SEQ ID NO:16:											
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 44 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 											
(ii) MOLECULE TYPE: protein											

	(xi)	SEQ	JENCI	E DES	SCRI	PTIO	N: SI	EQ II	ON C	:16:							
	Tyr 1	Asp	His	Asp	Tyr 5	Glu	Thr	Thr	Lys	Glu 10	Tyr	Thr	Leu	Arg	Ile 15	Arg	
	Ala	Gln	Asp	Gly 20	Gly	Arg	Thr	Pro	Leu 25	Ser	Asn	Val	Ser	Gly 30	Leu	Val	
	Thr	Val	Gln 35	Val	Leu	Asp	Ile	Asn 40	Asp	Asn	Ala	Pro					
(2)	INFO	TAMS	ON I	FOR S	SEQ I	D NO):17:	:									
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 129 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear																
	(ii)	MOLI	ECULI	E TYI	PE: 0	DNA											
	(xi)	SEQ	JENCI	E DES	SCRII	PTION	1: SE	EQ II	ONO:	17:							
GGG	GGTC	A T	racg?	AGGAC	AAC	CGGC	ATGT	TAG	AGATO	CGA (CGTGC	CAGG	CC AC	GAGA	CTAC	}	60
GAC	CTAACO	C A	ATTC	CAGC	CA	TGC	AAGG	TCA	CAGT	CAA (CTC	ATCG	C C	CAA	(GAT	A	120
ACG	CCCCZ	4															129
(2)	INFO	TAM	ON I	FOR S	SEQ :	D NO	:18	:									
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear																
	(ii)	MOLE	ECULE	TYI	PE: P	prote	∍in										
	(xi)	SEQU	JENCI	E DES	CRII	PTION	1: SI	EQ II	ONO:	:18:							
	Arg 1	Gly	Val	Asp	Tyr 5	Glu	Glu	Asn	Gly	Met 10	Leu	Glu	Ile	Asp	Val 15	Gln	
	Ala	Arg	Asp	Leu 20	Gly	Pro	Asn	Pro	Ile 25	Pro	Ala	His	Сув	Lys 30	Val	Thr	

Val Lys Leu Ile Asp Arg Asn Asp Asn Ala Pro 35

60

120

, <u> </u>	
(2) INFORMATION FOR SEQ ID NO:19:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:	
AAGGGGTTGG ACTACGAAGA CACCAAACTC CATGAGATTT ACATCCAGGC CAAAGACAAA	60
GGTGCCAATC CGGAAGGAGC GCATTGCAAA GTACTGGTAG AGGTTGTGGA CGTTAACGAC	120
AATGCCCCTC A	131
(2) INFORMATION FOR SEQ ID NO:20:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear (ii) MOLECULE TYPE: protein	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:	
Lys Gly Leu Asp Tyr Glu Asp Thr Lys Leu His Glu Ile Tyr Ile Gln 1 5 10 15	
Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys Lys Val Leu 20 25 30	
Val Glu Val Val Asp Val Asn Asp Asn Ala Pro 35 40	
(2) INFORMATION FOR SEQ ID NO:21:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

AAGGGTTTGG ACTTTGAGCA AGTAGATGTC TACAAAATCC GCGTTGACGC GACGGACAAA

GGACACCCTC CGATGGCAGG CCATTGCACT GTTTTAGTGA GGGTATTGGA TGAAAACGAC

AATGCGCCTC T	131
(2) INFORMATION FOR SEQ ID NO:22:	
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 	
(ii) MOLECULE TYPE: protein	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:	
Lys Gly Leu Asp Phe Glu Gln Val Asp Val Tyr Lys Ile Arg Val As	ip
Ala Thr Asp Lys Gly His Pro Pro Met Ala Gly His Cys Thr Val Le	eu
Val Arg Val Leu Asp Glu Asn Asp Asn Ala Pro 35 40	
(2) INFORMATION FOR SEQ ID NO:23:	
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 134 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 	
(ii) MOLECULE TYPE: cDNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:	
AAGGGTATAG ACTTCGAGCA GATCAAGGAC TTCAGCTTTC AAGTGGAAGC CCGGGACGCC	60
GGCAGTCCCC AGGCGCTGTC CGGCAACTGC ACTGTCAACA TCTTGATAGT GGATCAGAAC	120
GACAACGCCC CTAA	134
(2) INFORMATION FOR SEQ ID NO:24:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 44 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: protein	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:	
Lys Gly Ile Asp Phe Glu Gln Ile Lys Asp Phe Ser Phe Gln Val Gl	.u

Ala	Arg	Asp	Ala	Gly	Ser	Pro	Gln	Ala	Leu	Ala	Gly	Asn	Thr	Thr	Val
			20	_				25			•		30		

Asn Ile Leu Ile Val Asp Gln Asn Asp Asn Ala Pro 35 40

- (2) INFORMATION FOR SEQ ID NO:25:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 134 base pairs
 - (B) TYPE: nucleic acid (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:

AAGCCGTTCG ACTATGAGCA AACCGCCAAC ACGCTGGCAC AGATTGACGC CGTGCTGGAA

AAACAGGGCA GCAATAAATC GAGCATTCTG GATGCCACCA TTTTCCTGGC CGATAAAAAC

GACAATGCGC CAGA

134

- (2) INFORMATION FOR SEQ ID NO:26:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 44 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:

Lys Pro Phe Asp Tyr Glu Gln Thr Ala Asn Thr Leu Ala Gln Ile Asp 1 5 10 15

Ala Val Leu Glu Lys Gln Gly Ser Asn Lys Ser Ser Ile Leu Asp Ala 20 25 30

Thr Ile Phe Leu Ala Asp Lys Asn Asp Asn Ala Pro 35 40

- (2) INFORMATION FOR SEQ ID NO:27:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 131 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA

- 45 -	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:	
AAGCGGCTGG ATTTCGAACA GTTCCAGCAG CACAAGCTGC TCGTAAGGGC TGTTGATGGA	60
GGAATGCCGC CACTGAGCAG CGATGTGGTC GTCACTGTGG ATGTCACCGA CCTCAACGAT	120
AACGCGCCCT A	131
(2) INFORMATION FOR SEQ ID NO:28:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: protein	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:28:	
Lys Arg Leu Asp Phe Glu Gln Phe Gln Gln His Lys Leu Leu Val Arg 1 5 10 15	
Ala Val Asp Gly Gly Met Pro Pro Leu Ser Ser Asp Val Val Thr 20 25 30	
Val Asp Val Thr Asp Leu Asn Asp Asn Ala Pro 35 40	
(2) INFORMATION FOR SEQ ID NO:29:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:	
	60
	20
	31
(2) INFORMATION FOR SEQ ID NO:30:	
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 	
(ii) MOLECULE TYPE: protein	

	(xi)	SEQ	UENC	E DE	SCRI	PTIO	N: S	EQ I	D NO	:30:							
	Lys 1	Gly	Ile	Asp	Phe 5	Glu	Ser	Glu	Asn	Tyr 10	Tyr	Glu	Phe	Asp	Val 15	Arg	
	Ala	Arg	Asp	Gly 20	Gly	Ser	Pro	Ala	Met 25	Glu	Gln	His	Сув	Ser 30	Leu	Arg	
	Val	Asp	Leu 35	Leu	Asp	Val	Asn	Asp 40	Asn	Ala	Pro						
(2)	INFO	RMAT:	ION I	FOR S	SEQ 1	D NO	31:	:									
	(i)	(B)	JENCI LEI TYI STI	IGTH: PE: 1 RANDI	: 131 nucle EDNES	l bas eic a SS: s	se pa acid singl	airs									
	(ii)	MOLI	CULE	TYE	PE: c	DNA											
٠	(xi)	SEQU	JENCE	DES	CRIE	PTION	: SE	Q II	NO:	31:							
AAG	GCATTO	G AC	TTTG	AGGC	cce	GCGA	CTG	TAT	rcger	GA C	CAGTI	CAGG	C C	ACGG2	ACCG	.	60
GGC	STGCCC	T C	CTCA	CCGG	GCG	TGCC	GAA	GCG	CTTAT	CC F	GCT	CTAG	A TO	TCA	ACGAC	:	120
AAC	GCACCO	A T															131
(2)	INFOR	ITAM	ON F	OR S	EQ I	D NO	:32:										
	(i)	(B)	ENCE LEN TYP STR TOP	GTH: E: a ANDE	43 mino DNES	amin aci S: s	o ac d ingl	ids									
	(ii)	MOLE	CULE	TYP	E: p	rote	in										
	(xi)	SEQU	ENCE	DES	CRIP	TION	: SE	Q II	NO:	32:							
	Lys 1	Ala	Leu	Asp	Phe 5	Glu	Ala	Arg	Arg	Leu 10	Tyr	Ser	Leu	Thr	Val 15	Gln	
	Ala	Thr	Asp .	Arg 20	Gly	Val	Pro	Ser	Leu 25	Thr	Gly	Arg	Ala	Glu	Ala	Leu	

Ile Gln Leu Leu Asp Val Asn Asp Asn Ala Pro 35 40

60

120

(2) INFORMATION FOR SEQ ID NO:33:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 125 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:33:	
AAGCCAATTG ATTACGAGGC AACTCCATAC TATAACATGG AAATTGTAGC CACAGACAGC	60
GGAGGTCTTT CGGGAAAATG CACTGTGTCT ATACAGGTGG TGGATGTGAA CGACAACGCC	120
CCCAA	125
(2) INFORMATION FOR SEQ ID NO:34:	
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 41 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 	
(ii) MOLECULE TYPE: protein	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:34:	
Lys Pro Ile Asp Tyr Glu Ala Thr Pro Tyr Tyr Asn Met Glu Ile Val 1 5 10 15	
Ala Thr Asp Ser Gly Gly Leu Ser Gly Lys Cys Thr Val Ser Ile Gln 20 25 30	
Val Val Asp Val Asn Asp Asn Ala Pro 35 40	
(2) INFORMATION FOR SEQ ID NO:35:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 446 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:35:

AAGCGGGTAG ACTTCGAAAT GTGCAAAAGA TTTTACCTTG TGGTGGAAGC TAAAGACGGA

GGCACCCCAG CCCTCAGCAC GGCAGCCACT GTCAGCATCG ACCTCACAGA TGTGAATGAT

AACCCTCCTC	GGTTCAGCCA	AGATGTCTAC	AGTGCTGTCA	TCAGTGAGGA	TGCCTTAGAG	180
GGGGACTCTG	TCATTCTGCT	GATAGCAGAA	GATGTGGATA	GCAAGCCTAA	TGGACAGATT	240
CGGTTTTCCA	TCGTGGGTGG	AGATAGGGAC	AATGAATTTG	CTGTCGATCC	AATCTTGGGA	300
CTTGTGAAAG	TTAAGAAGAA	ACTGGACCGG	GAGCGGGTGT	CAGGATACTC	CCTGCTCATC	360
CAGGCAGTAG	ATAGTGGCAT	TCCTGCAATG	TCCTCAACGA	CAACTGTCAA	CATTGATATT	420
TCTGATGTGA	ACGACAACGC	CCCCT				446

- (2) INFORMATION FOR SEQ ID NO:36:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 148 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:
 - Lys Arg Val Asp Phe Glu Met Cys Lys Arg Phe Tyr Leu Val Val Glu 1 5 10 15
 - Ala Lys Asp Gly Gly Thr Pro Ala Leu Ser Thr Ala Ala Thr Val Ser 20 25 30
 - Ile Asp Leu Thr Asp Val Asn Asp Asn Pro Pro Arg Phe Ser Gln Asp
 - Val Tyr Asp Ala Val Ile Ser Glu Asp Ala Leu Glu Gly Asp Ser Val 50 60
 - Ile Leu Leu Ile Ala Glu Asp Val Asp Ser Lys Pro Asn Gly Gln Ile 65 70 75 80
 - Arg Phe Ser Ile Val Gly Gly Asp Arg Asp Asn Glu Phe Ala Val Asp 85 90 95
 - Pro Ile Leu Gly Leu Val Lys Val Lys Lys Leu Asp Arg Glu Arg
 100 105 110
 - Val Ser Gly Tyr Ser Leu Leu Ile Gln Ala Val Asp Ser Gly Ile Pro 115 120 125
 - Ala Met Ser Ser Thr Thr Val Asn Ile Asp Ile Ser Asp Val Asn 130 135 140

Asp Asn Ala Pro 145

(2) INFORMATION FOR SEQ ID NO:37:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 440 base pairs(B) TYPE: nucleic acid

 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:

AAGGGGGTTG	ATTATGAGAC	AAACCCACGG	CTACGACTGG	TGCTACAGGC	AGAGAGTGGA	60
GGAGCCTTTG	CTTTCTCGGT	GCTGACCCTG	ACCCTTCAAG	ATGCCAATGA	CAATGCTCCC	120
CGTTTCCTGC	AGCCTCACTA	CGTGGCTTTC	CTGCCAGAGT	CCCGACCCTT	GGAAGGGCCC	180
CTGCTGCAGG	TGGAAGCAGA	CGACCTGGAT	CAAGGCTCTG	GAGGACAGAT	CTCCTACAGT	240
CTGGCTGCAT	CCCAGCCAGC	ACGGGGCTTG	TTCCATGTAG	ACCCAGCCAC	AGGCACTATC	300
ACTACCACAG	CCATCCTGGA	CCGGGAAATC	TGGGCTGAAA	CACGGCTGGT	ACTGATGGCC	360
ACAGACAGAG	GAAGCCCAGC	ATTGGTGGGC	TCAGCTACCC	TGACAGTGAT	GGTCATCGAT	420
ACCAACGACA	ATGCTCCCCT					440

(2) INFORMATION FOR SEQ ID NO:38:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 146 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:38:

- Lys Gly Val Asp Tyr Glu Thr Asn Pro Arg Leu Arg Leu Val Leu Gln 1 15
- Ala Glu Ser Gly Gly Ala Phe Ala Phe Ser Val Leu Thr Leu
- Gln Asp Ala Asn Asp Asn Ala Pro Arg Phe Leu Gln Pro His Tyr Val
- Ala Phe Leu Pro Glu Ser Arg Pro Leu Glu Gly Pro Leu Leu Gln Val 50
- Glu Ala Asn Asp Leu Asp Gln Gly Ser Gly Gly Gln Ile Ser Tyr Ser 65 70 75 80
- Leu Ala Ala Ser Gln Pro Ala Arg Gly Leu Phe His Val Asp Pro Ala

Thr	Gly	Thr	Ile	Thr	Thr	Thr	Ala	Ile	Leu	Asp	Arg	Glu	Ile	Trp	Ala
			100					105					110		

Glu Thr Arg Leu Val Leu Met Ala Thr Asp Arg Gly Ser Pro Ala Leu 115 120 125

Val Gly Ser Ala Thr Leu Thr Val Met Val Ile Asp Thr Asn Asp Asn 130 135 140

Ala Pro 145

- (2) INFORMATION FOR SEQ ID NO:39:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 124 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: CDNA
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:39:

AAGGTCTCGA TTATGAGGCA ACTCCATATT ATAACGTGGA AATTGTAGCC ACAGATGGTG 60
GGGGCCTTTC AGGAAAATGC ACTGTGGCTA TAGAAGTGGT GGATGTGAAC GACGGCGCTC 120
CAAT

- (2) INFORMATION FOR SEQ ID NO:40:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 41 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:40:

Lys Gly Leu Asp Tyr Glu Ala Thr Pro Tyr Tyr Asn Val Glu Ile Val 1 5 10 15

Ala Thr Asp Gly Gly Ala Phe Asp Glu Asn Cys Thr Val Ala Ile Glu 20 25 30

Val Val Asp Val Asn Asp Asn Ala Pro 35 40

- (2) INFORMATION FOR SEQ ID NO:41:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 8 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:41:

Asp Xaa Asn Glu Xaa Pro Xaa Phe

- (2) INFORMATION FOR SEQ ID NO:42:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 8 amino acids

 - (B) TYPE: amino acid (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:42:

Asp Xaa Asp Glu Xaa Pro Xaa Phe

- (2) INFORMATION FOR SEQ ID NO:43:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 9 amino acids(B) TYPE: amino acid

 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: DNA
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:43:

Asp Xaa Asn Asp Asn Xaa Pro Xaa Phe

- (2) INFORMATION FOR SEQ ID NO:44:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 131 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:44:	
AAGCGGATGG ATTTTGAAGA CACCAAACTC CATGAGATTT ACATCCAGGC CAAAGACAAA	50
GGTGCCAATC CCGAAGGAGC GCATTGCAAA GTACTTGTAG AGGTTGTAGA CGTAAACGAC 12	2 C
AACGCCCCAG T	31
(2) INFORMATION FOR SEQ ID NO:45:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: protein	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:45:	
Leu Arg Met Asp Phe Glu Asp Thr Lys Leu His Glu Ile Tyr Ile Gln 1 5 10 15	
Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys Lys Val Leu 20 25 30	
Val Glu Val Val Asp Val Asn Asp Asn Ala Pro 35 40	
(2) INFORMATION FOR SEQ ID NO:46:	
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 	
(ii) MOLECULE TYPE: cDNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:46:	
AAGGCTTTGG ATTACGAGGA TCAGAGAGAG TTCCAACTAA CAGCTCATAT AAACGACGGA 6	0
GGTACCCCAG TCTTAGCCAC CAACATCAGC GTGAACGTAT TTGTTACTGA CCGCAATGAT 12	0
AACGCCCCCT A	1
(2) INFORMATION FOR SEQ ID NO:47:	
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 	
(ii) MOLECULE TYPE: protein	

	(xi)	SEQ	JENC	E DE	SCRI	PTIO	N: SI	EQ II	ON C	:47:							
	Lys 1	Ala	Leu	Asp	Tyr 5	Glu	Asp	Gln	Arg	Glu 10	Phe	Gln	Leu	Thr	Ala 15	His	
	Ile	Asn	Asp	Gly 20	Gly	Thr	Pro	Val	Leu 25	Ala	Thr	Asn	Ile	Ser 30	Val	Asn	
	Val	Phe	Val 35	Thr	Asp	Arg	Asn	Asp 40	Asn	Ala	Pro						
(2)	INFO	TAMS	ON I	FOR S	SEQ 1	D NO	:48	:									
	(i)	(A) (B) (C)	LEI TYI STI	NGTH: PE: 1 RANDE	ARACT : 131 nucle EDNES GY: 1	l bas eic a SS: s	se pa acid sing]	airs									
	(ii)	MOLE	CULE	TYI	PE: c	DNA											
	(xi)	SEQU	ENCE	E DES	CRIE	PTION	l: SE	EQ II	NO:	48:							
AAGC	GCTTC	G AC	TAC	AGG	A GAG	TAAC	TAA	TATO	RAAT	TC F	ACGTO	GATO	C TA	CAGA	TAAF	4	60
GGAT	ACCC	C CI	ATGO	TTGC	TCA	CTGC	ACC	GTAC	CTCGI	GG G	BAATO	TTGG	A TO	AAAA	TGAC	3	120
AACG	CACCO	A T															131
(2)	INFOF	ITAM	ON F	OR S	EQ I	D NC	:49:										
	(i)	(A) (B) (C)	LEN TYP STF	IGTH: PE: a NANDE	ARACT 43 uminc DNES Y: 1	amin aci S: s	o ac d ingl	ids									
	(ii)	MOLE	CULE	TYP	E: p	rote	in										
	(xi)	SEQU	ENCE	DES	CRIP	TION	: SE	Q ID	NO:	49:							
	Lys 1	Arg	Leu	Asp	Tyr 5	Glu	Glu	Ser	Asn	Asn 10	Tyr	Glu	Ile	His	Val 15	Asp	
	Ala	Thr	Asp	Lys 20	Gly	Tyr	Pro	Pro	Met 25	Val	Ala	His	Сув	Thr 30	Val	Leu	

Val Gly Ile Leu Asp Glu Asn Asp Asn Ala Pro 35 40

60

120

- 34 -	
(2) INFORMATION FOR SEQ ID NO:50:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:50:	
AAACCGGTGG ACTACGAGAA AGTCAAAGAC TATACCATCG AGATCGTGGC TGTGGATTCC	60
GGCAACCCTC CACTCTCTAG CACCAACTCC CTCAAGGTGC AGGTGGTAGA CGTCAACGAT	120
AACGCCCCTC T	131
(2) INFORMATION FOR SEQ ID NO:51: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear (ii) MOLECULE TYPE: protein	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:51:	
Lys Pro Val Asp Tyr Glu Lys Val Lys Asp Tyr Thr Ile Glu Ile Val	
Ala Val Asp Ser Gly Asn Pro Pro Leu Ser Ser Thr Asn Ser Leu Lys 20 25 30	
Val Gln Val Val Asp Val Asn Asp Asn Ala Pro 35 40	
(2) INFORMATION FOR SEQ ID NO:52:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:52:

AAGCCTTTTG ATTTCGAGGA CACCAAACTC CATGAGATTT ACATCCAGGC CAAAGACAAG

GGCGCCAATC CCGAAGGAGC ACATTGCAAA GTGTTGGTGG AGGTTGTGGA TGTGAACGAC

AAT	GCCCCTC A	131
(2)	INFORMATION FOR SEQ ID NO:53:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
	(ii) MOLECULE TYPE: protein	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO:53:	
	Lys Pro Phe Asp Phe Glu Asp Thr Lys Leu His Glu Ile Tyr Ile Gln 1 5 10 15	
	Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys Lys Val Leu 20 25 30	
	Val Glu Val Val Asp Val Asn Asp Asn Ala Pro 35 40	
(2)	INFORMATION FOR SEQ ID NO:54:	
	 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 122 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 	
	(ii) MOLECULE TYPE: cDNA	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO:54:	
AAAG	GGTGTCG ATTACGAGGT GAGTCCACGG CTGCGACTGG TGCTGCAGGC AGAGAGTCGA	60
GGAG	SCCTTTG CCTTCACTGT GCTGACCCTG ACCCTGCAAG ATGCCAACGA CAACGCCCCG	120
AG		122
(2)	INFORMATION FOR SEQ ID NO:55:	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 40 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
	(ii) MOLECULE TYPE: protein	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO:55:	

Lys Gly Val Asp Tyr Glu Val Ser Pro Arg Leu Arg Leu Val Leu Gln 1 5 10 15

Ala	Glu	Ser	Arg	Gly	Ala	Phe	Ala	Phe	Thr	Val	Leu	Thr	Leu	Thr	Leu
			20	_				25					30		

Gln Asp Ala Asn Asp Asn Ala Pro 35 40

- (2) INFORMATION FOR SEQ ID NO:56:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid

 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:56:

AAAGGGATTG ATTACGAGCA GTTGAGAGAC CTACAGCTGT GGGTGACAGC CAGCGACAGC 60 GGGGACCCGC CTCTTAGCAG CAACGTGTCA CTGAGCCTGT TTGTGCTGGA CCAGAACGAC 120 AACGCCCCC T 131

- (2) INFORMATION FOR SEQ ID NO:57:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 43 amino acids

 - (B) TYPE: amino acid (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:57:

Lys Gly Ile Asp Tyr Glu Gln Leu Arg Asp Leu Gln Leu Trp Val Thr

Ala Ser Asp Ser Gly Asp Pro Pro Leu Ser Ser Asn Val Ser Leu Ser

Leu Phe Val Leu Asp Gln Asn Asp Asn Ala Pro 35 40

- (2) INFORMATION FOR SEQ ID NO:58:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 125 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA

- 3/ -	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:58:	
AAGGCGGTCG ATTTTGAGCG CACATCCTCT TATCAACTCA TCATTCAGGC CACCAATATG	60
GCAGGAATGG CTTCCAATGC TACAGTCAAT ATTCAGATTG TTGATGAAAA CGACAACGCC 1	20
CCCCA	25
(2) INFORMATION FOR SEQ ID NO:59:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 41 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: protein	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:59:	
Lys Ala Val Asp Phe Glu Arg Thr Ser Ser Tyr Gln Leu Ile Ile Gln 1 5 10 15	
Ala Thr Asn Met Ala Gly Met Ala Ser Asn Ala Thr Val Asn Ile Gln 20 25 30	
Ile Val Asp Glu Asn Asp Asn Ala Pro 35 40	
(2) INFORMATION FOR SEQ ID NO:60:	
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 	
(ii) MOLECULE TYPE: cDNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:60:	
AAACGGCTAG ACTTTGAAAA GATACAAAAA TATGTTGTAT GGATAGAGGC CAGAGATGGT	60
GGTTTCCCTC CTTTCTCCTC TTACGAGAAA CTTGATATAA CAGTATTAGA TGTCAACGAT 1:	20
AACGCGCCTA A	31
(2) INFORMATION FOR SEQ ID NO:61:	
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 	

(ii) MOLECULE TYPE: protein

(xi)	SEQ	JENC	E DE	SCRI	PTIO	N: S	EQ I	on o	:61:						
Lys 1	Arg	Leu	Asp	Phe 5	Glu	Lys	Ile	Gln	Lys 10	Tyr	Val	Val	Trp	Ile 15	Glu
Ala	Arg	Asp	Gly 20	Gly	Phe	Pro	Pro	Phe 25	Ser	Ser	Tyr	Glu	Lys 30	Leu	Asp
Ile	Thr	Val 35	Leu	Asp	Val	Asn	Asp 40	Asn	Ala	Pro					
NFOE	TAMS	ON I	FOR S	SEQ 1	D NO): 62:	:								
(i)	(A)	LEN	CHA	: 131	bas	se pa									

- (2) I

 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:62:

AAGGGGATCG ATTATGAGAA GGTCAAAGAC TACACCATTG AGATTGTGGC TGTGGACTCT 60 GGCAACCCCC CACTCCCAG CACTAACTCC CTCAAGGTGC AGGTGGTGGA CGTCAATGAC 120 AACGCACCGT G 131

- (2) INFORMATION FOR SEQ ID NO:63:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 43 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:63:
 - Lys Gly Ile Asp Tyr Glu Lys Val Lys Asp Tyr Thr Ile Glu Ile Val
 - Ala Val Asp Ser Gly Asn Pro Pro Leu Ser Ser Thr Asn Ser Leu Lys
 - Val Gln Val Val Asp Val Asn Asp Asn Ala Pro 40

(2) INFORMATION FOR SEQ ID NO:64:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:64:	
AAGGGACTCG ACTACGAGGA TCGGCGGGAA TTTGAATTAA CAGCTCATAT CAGCGATGGG	60
GGCACCCCGG TCCTAGCCAC CAACATCAGC GTGAACATAT TTGTCACTGA TCGCAACGAT	120
AATGCCCCCG T	131
(2) INFORMATION FOR SEQ ID NO:65:	
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 	
(ii) MOLECULE TYPE: protein	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:65:	
Lys Gly Leu Asp Tyr Glu Asp Arg Arg Glu Phe Glu Leu Thr Ala His 1 5 10 15	
Ile Ser Asp Gly Gly Thr Pro Val Leu Ala Thr Asn Ile Ser Val Asn 20 25 30	
Ile Phe Val Thr Asp Arg Asn Asp Asn Ala Pro 35 40	
(2) INFORMATION FOR SEQ ID NO:66:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 470 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:66:

AAGGGTTTGG ACTACGAGAC CACACAGGCC TACCAGCTCA CGGTCAACGC CACAGATCAA

GACAACACCA GGCCTCTGTC CACCCTGGCC AACTTGGCCA TCATCATCAC AGATGTCCAG

60

120

GACATGGACC	CCATCTTCAT	CAACCTGCCT	TACAGCACCA	ACATCTACGA	GCATTCTCCT	180
CCGGGCACGA	CGGTGCGCAT	CATCACCGCC	ATAGACCAGG	ATCAAGGACG	TCCCCGGGC	240
ATTGGCTACA	CCATCGTTTC	AGGGAATACC	AACAGCATCT	TTGCCCTGGA	CTACATCAGC	300
GGAGTGCTGA	CCTTGAATGG	CCTGCTGGAC	CGGGAGAACC	CCCTGTACAG	CCATGGCTTC	360
ATCCTGACTG	TGAAGGCAC	GGAGCTGAAC	GATGACCGCA	CCCCATCTGA	CGCTACAGTC	420
ACCACGACCT	TCAATATCCT	GGTTATTGAC	ATCAACGACA	ACGCCCCACT		470

- (2) INFORMATION FOR SEQ ID NO:67:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 156 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:67:
 - Lys Gly Leu Asp Tyr Glu Thr Thr Gln Ala Tyr Gln Leu Thr Val Asn 1 5 10 15
 - Ala Thr Asp Gln Asp Asn Thr Arg Pro Leu Ser Thr Leu Ala Asn Leu 20 25 30
 - Ala Ile Ile Ile Thr Asp Val Gln Asp Met Asp Pro Ile Phe Ile Asn 35 40 45
 - Leu Pro Tyr Ser Thr Asn Ile Tyr Glu His Ser Pro Pro Gly Thr Thr 50 60
 - Val Arg Ile Ile Thr Ala Ile Asp Gln Asp Gln Gly Arg Pro Arg Gly 65 70 75 80
 - Ile Gly Tyr Thr Ile Val Ser Gly Asn Thr Asn Ser Ile Phe Ala Leu 85 90 95
 - Asp Tyr Ile Ser Gly Val Leu Thr Leu Asn Gly Leu Leu Asp Arg Glu 100 105 110
 - Asn Pro Leu Tyr Ser Gly Gly Phe Ile Leu Thr Val Lys Gly Thr Glu 115 120 125
 - Leu Asn Asp Asp Arg Thr Pro Ser Asp Ala Thr Val Thr Thr Thr Phe 130 135
 - Asn Ile Leu Val Ile Asp Ile Asn Asp Asn Ala Pro 145 150 155

- 61 -	
(2) INFORMATION FOR SEQ ID NO:68:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:68:	
AAGGGGGTCG ATTACGAGGT ACTACAGGCC TTTGAGTTCC ACGTGAGCGC CACAGACCGA	60
GGCTCACCGG GGCTCAGCAG CCAGGCTCTG GTGCGCGTGG TGGTGCTGGA CGACAATGAC	120
AACGCTCCCG T	L31
(2) INFORMATION FOR SEQ ID NO:69:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: protein	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:69:	
Lys Gly Val Asp Tyr Glu Val Leu Gln Ala Phe Glu Phe His Val Ser 1 5 10 15	
Ala Thr Asp Arg Gly Ser Pro Gly Leu Ser Ser Gln Ala Leu Val Arg 20 25 30	
Val Val Leu Asp Asp Asn Asp Asn Ala Pro 35 40	
(2) INFORMATION FOR SEQ ID NO:70:	
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear	
(ii) MOLECULE TYPE: cDNA	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:70:	

AAGGGGCTGG ATTATGAGCA GTTCCAGACC CTACAACTGG GAGTGACCGC TAGTGACAGT

GGAAACCCAC CATTAAGAAG CAATATTTCA CTGACCCTTT TCGTGCTGGA CCAGAATGAT

60

120

	GCCCC			FOR	CEO :	TO W	0.71	_									131
(2)	INFORMATION FOR SEQ ID NO:71:																
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear																
	(ii)	MOL	ECULI	E TY	PE:]	prot	ein										
	(xi)	SEQ	UENCI	E DE	SCRI	PTIO	N: S	EQ I	D NO	:71:							
	Lys 1	Gly	Leu	Asp	Tyr 5	Glu	Gln	Phe	Gln	Thr 10	Leu	Gln	Leu	Gly	Val 15	Thr	
	Ala	Ser	Asp	Ser 20	Gly	Asn	Pro	Pro	Leu 25	Arg	Ser	Asn	Ile	Ser 30	Leu	Thr	
	Leu	Phe	Val 35	Leu	Asp	Gln	Asn	Asp 40	Asn	Ala	Pro						
(2)	INFO	RMAT:	ION I	FOR :	SEQ :	ID NO): 72	:									
	(i)	(A (B (C	UENCI) LEI) TYI) STI) TOI	NGTH PE: : RAND	: 13: nucle EDNE:	l bas eic a SS: s	se pa acid sing	airs									
	(ii)	MOLI	ECULE	E TY	PE: (CDNA											
	(xi)	SEO	JENCE	E DES	SCRII	PTIO	V: 51	EO TI	סא מ	.72.							
AAGC	GGGT										CAT'	raag:	GC C	CAGG	ATGG	G	60
	cccc																120
AATG	cccc	GG A															131
(2)	INFO	RMAT	ON E	FOR S	SEQ 1	ID NO):73:	:									
	(i)	(A) (B) (C)	JENCE LEN TYP STF	NGTH PE: 8 RANDI	: 43 emino EDNES	amin caci	no ad id sing:	cids									
	(ii)	MOLE	ECULE	TYI	PE: p	pepti	ide										

Lys Arg Val Asp Tyr Glu Asp Val Gln Lys Tyr Ser Leu Ser Ile Lys 1 5 10 15

CCGAT

Ala Gln Asp Gly Arg Pro Pro Leu Ile Asn Ser Ser Gly Val Val Ser 30

Val Gln Val Leu Asp Val Asn Asp Asn Ala Pro 35 40

- (2) INFORMATION FOR SEQ ID NO:74:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 125 base pairs (B) TYPE: nucleic acid

 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:74: AAACCGGTAG ACTTTGAGCT ACAGCAGTTC TATGAAGTAG CTGTGGTGGC TTGGAACTCT 60 GAGGGATTTC ATGTCAAAAG GGTCATTAAA GTGCAACTTT TAGATGACAA CGACAATGCC 120

125

- (2) INFORMATION FOR SEQ ID NO:75:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 41 amino acids

 - (B) TYPE: amino acid (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:75:

Lys Pro Val Asp Phe Glu Leu Gln Gln Phe Tyr Glu Val Ala Val Val

Ala Trp Asn Ser Glu Gly Phe His Val Lys Arg Val Ile Lys Val Gln

Leu Leu Asp Asp Asn Asp Asn Ala Pro 35 40

- (2) INFORMATION FOR SEQ ID NO:76:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 125 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA

√
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:76:
AAGGGATTAG ATTTTGAAAC TTTGCCCATT TACACATGA TAATACAAGG AACTAACATG 60
GCTGGTTTGT CCACTAATAC AACGGTTCTA GTTCACTTGC AGGATGAGAA TGATAACGCC 120
CCAAA 129
(2) INFORMATION FOR SEQ ID NO:77:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 41 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear
(ii) MOLECULE TYPE: protein
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:77:
Lys Gly Leu Asp Phe Glu Thr Leu Pro Ile Tyr Thr Leu Ile Ile Gln 1 5 10 15
Gly Thr Asn Met Ala Gly Leu Ser Thr Asn Thr Thr Val Leu Val His 20 25 30
Leu Gln Asp Glu Asn Asp Asn Ala Pro 35 40
(2) INFORMATION FOR SEQ ID NO:78:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 134 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear
(ii) MOLECULE TYPE: cDNA
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:78:
AAGCGGGCGG ATTTCGAGGC GATCCGGGAG TACAGTCTGA GGATCAAAGC GCAGGACGGG 60
GGGCGGCCTC CCCTCAGCAA CACCACGGGC ATGGTCACAG TGCAGGTCGT GGACGTCAAT 120
GACAACGCAC CCCT 134
(2) INFORMATION FOR SEQ ID NO:79:
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 44 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(2)

(xi)	SEQU	JENCI	E DES	CRI	PTIO	1: SI	II Q	ON C	79:						
Lys 1	Arg	Ala	Asp	Phe 5	Glu	Ala	Ile	Arg	Glu 10	Tyr	Ser	Leu	Arg	Ile 15	Lys
Ala	Gln	Asp	Gly 20	Gly	Arg	Pro	Pro	Leu 25	Ser	Asn	Thr	Thr	Gly 30	Met	Val
Thr	Val	Gln 35	Val	Val	Asp	Val	Asn 40	Asp	Asn	Ala	Pro				
INFOR	CTAMS	ION I	FOR S	SEQ :	ID NO	0:80	:								
(i)	(A) (B) (C)	LEI TYI STI	NGTH PE: 1 RANDI	: 13: nucle EDNE:	reris l bas eic s ss: s lines	se pa acid sing:	airs								
(ii)	MOLI	ECULI	E TY	PE:	CDNA										

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:80:

AAGCGGTTGG ATTACGAAAA GGCATCGGAA TATGAAATCT ATGTTCAAGC CGCTGACAAA 60 GGCGCTGTCC CTATGGCTGG CCATTGCAAA GTGTTGCTGG AGATCGTGGA TGTCAACGAC 120 131 AACGCCCCCT T

- (2) INFORMATION FOR SEQ ID NO:81:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 43 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:81:

Lys Arg Leu Asp Tyr Glu Lys Ala Ser Glu Tyr Glu Ile Tyr Val Gln

Ala Ala Asp Lys Gly Ala Val Pro Met Ala Gly His Cys Lys Val Leu 20 25 30

Leu Glu Ile Val Asp Val Asn Asp Asn Ala Pro

- 00 -										
(2) INFORMATION FOR SEQ ID NO:82:										
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear										
(ii) MOLECULE TYPE: cDNA										
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:82:										
AAGGGGATCG ATTATGAGGA TCAGGTCTCT TACACATTAG CAGTAACAGC ACATGACTAT	60									
GGCATCCCTC AAAAATCAGA CACTACCTAT TTGGAAATCT TAGTAATTGA TGTTAACGAC	120									
AACGCGCCCC A	131									
(2) INFORMATION FOR SEQ ID NO:83:										
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear										
(ii) MOLECULE TYPE: protein										
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:83:										
Lys Gly Ile Asp Tyr Glu Asp Gln Val Ser Tyr Thr Leu Ala Val Thr 1 5 10 15										
Ala His Asp Tyr Gly Ile Pro Gln Lys Ser Asp Thr Thr Tyr Leu Glu 20 25 30										
Ile Leu Val Ile Asp Val Asn Asp Asn Ala Pro 35 40										
(2) INFORMATION FOR SEQ ID NO:84:										
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 										
(ii) MOLECULE TYPE: cDNA										

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:84:

AAAGGGTTAG ATTTCGAGGG CACTAAAGAT TCAGCGTTTA AAATAGTGGC AGCTGACACA

GGGAAGCCCA GCCTCAACCA GACAGCCCTG GTGAGAGTAG AGCTGGAGGA TGAGAACGAC

60

120

AACGCCCCAA T	131											
(2) INFORMATION FOR SEQ ID NO:85:												
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 												
(ii) MOLECULE TYPE: protein												
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:85:												
Lys Gly Leu Asp Phe Glu Gly Thr Lys Asp Ser Ala Phe Lys Ile Val												
Ala Ala Asp Thr Gly Lys Pro Ser Leu Asn Gln Thr Ala Leu Val Arg 20 25 30												
Val Glu Leu Glu Asp Glu Asn Asp Asn Ala Pro 35 40												
(2) INFORMATION FOR SEQ ID NO:86:												
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 130 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear												
(ii) MOLECULE TYPE: cDNA												
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:86:												
AAGGGTGTGG ATTTTGAAAG TGTGCGTAGC TACAGGCTGG TTATTCGTGC TCAAGATGGA	60											
GGCAGCCCCT CCAGAAGTAA CACCACCCAG CTCTTGGTCA ACGTCATCGA TCGAATGACA	120											
ATGCGCCGCT	130											
(2) INFORMATION FOR SEQ ID NO:87:												
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear												
(ii) MOLECULE TYPE: protein												

Lys Gly Val Asp Phe Glu Ser Val Arg Ser Tyr Arg Leu Val Ile Arg 1 5 10 15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:87:

Ala	Gln	Asp	Gly	Gly	Ser	Pro	Ser	Arg	Ser	Asn	Thr	Thr	Gln	Leu	Lev
		_	20	-				25					30		

Val Asn Val Ile Asp Val Asn Asp Asn Ala Pro 35

- (2) INFORMATION FOR SEQ ID NO:88:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid

 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:88:

AAGGGTGTGG ACTTCGAGCT GACACATCTG TATGAGATTT GGATTGAGGC TGCCGATGGA 60 GACACGCCAA GTCTGCGTAG TGTAACTCTT ATAACGCTCA ACGTAACGGA TGCCAATGAC AATGCTCCCA A 131

- (2) INFORMATION FOR SEQ ID NO:89:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 43 amino acids

 - (B) TYPE: amino acid
 (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:89:

Lys Gly Val Asp Phe Glu Leu Thr His Leu Tyr Glu Ile Trp Ile Glu

Ala Ala Asp Gly Asp Thr Pro Ser Leu Arg Ser Val Thr Leu Ile Thr

Leu Asn Val Thr Asp Ala Asn Asp Asn Ala Pro 35

- (2) INFORMATION FOR SEQ ID NO:90:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 441 base pairs
 - (B) TYPE: nucleic acid (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA

	(x1) S	EQUENCE DESC	ERIPTION: SI	SÕ ID MO: AO:			
CAA	GGCGTTT	GATTTTGAAG	AGACAAGTAG	ATATGTGTTG	AGTGTGGAAG	CTAAGGATGG	60
AGG	AGTACAC	ACAGCTCACT	GTAATGTTCA	AATAGAAATT	GTTGACGAGA	ATGACAATGC	120
ccc	AGAGGTG	ACATTCATGT	CCTTCTCTAA	CCAGATTCCA	GAGGATTCAG	ACCTTGGAAC	180
TGT	AATAGCC	CTCATAAAAG	TGCGAGACAA	GGATTCTGGG	CAAAATGGCA	TGGTGACATG	240
CTA	TACTCAG	GAAGAAGTTC	CTTTCAAATT	AGAATCCACC	TCGAAGAATT	ATTACAAGCT	300
GGT	GATTGCT	GGAGCCCTAA	ACCGGGAGCA	GACAGCAGAC	TACAACGTCA	CAATCATAGC	360
CAC	CGACAAG	GGCAAACCAG	CCCTTTCCTC	CAGGACAAGC	ATCACCCTGC	ACATCTCCGA	420
CAT	CAACGAT	AATGCCCCCG	T				441

(2) INFORMATION FOR SEQ ID NO:91:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 146 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: protein
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:91:
- Lys Ala Phe Asp Phe Glu Glu Thr Ser Arg Tyr Val Leu Ser Val Glu
- Ala Lys Asp Gly Gly Val His Thr Ala His Cys Asn Val Gln Ile Glu 20 25 30
- Ile Val Asp Glu Asn Asp Asn Ala Pro Glu Val Thr Phe Met Ser Phe 35 40 45
- Ser Asn Gln Ile Pro Glu Asp Ser Asp Leu Gly Thr Val Ile Ala Leu 50 55
- Ile Lys Val Arg Asp Lys Asp Ser Gly Gln Asn Gly Met Val Thr Cys 65 70 75 80
- Tyr Thr Gln Glu Glu Val Pro Phe Lys Leu Glu Ser Thr Ser Lys Asn 85 90 95
- Tyr Tyr Lys Leu Val Ile Ala Gly Ala Leu Asn Arg Glu Gln Thr Ala 100 105 110
- Asp Tyr Asn Val Thr Ile Ile Ala Thr Asp Lys Gly Lys Pro Ala Leu 115 120 125
- Ser Ser Arg Thr Ser Ile Thr Leu His Ile Ser Asp Ile Asn Asp Asn 130 135 140

Ala Pro

145

60 120 131

(2) INFORMATION FOR SEQ ID NO:92:													
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 131 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 													
(ii) MOLECULE TYPE: cDNA													
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:92:													
AAGCGAGTGG ATTACGAGGC CACTCGGAAT TATAAGCTGA GAGTTAAGGC TACTGATCTT													
GGGATTCCAC CGAGATCTTC TAACATGACA CTGTTCATTC ATGTCCTTGA TGTTAACGAC													
AACGCTCCCT T													
(2) INFORMATION FOR SEQ ID NO:93:													
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids (B) TYPE: amino acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 													
(ii) MOLECULE TYPE: protein													
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:93:													
Lys Arg Val Asp Tyr Glu Ala Thr Arg Asn Tyr Lys Leu Arg Val Lys 1 10 15													
Ala Thr Asp Leu Gly Ile Pro Pro Arg Ser Ser Asn Met Thr Leu Phe 20 25 30													
Ile His Val Leu Asp Val Asn Asp Asn Ala Pro 35 40													
(2) INFORMATION FOR SEQ ID NO:94:													
 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 4104 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: single (D) TOPOLOGY: linear 													

(ii) MOLECULE TYPE: cDNA

(A) NAME/KEY: CDS (B) LOCATION: 495..3572

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:94:

(ix) FEATURE:

CCTCTATTCG ACA	ATTCTCTT TGGATTG1	TTT TGCTATAACT	TGAAATTTGG GATGTCAC	AA 60
ACGAAACTGT CAT	CTGTTTC CGCCAAAC	CTG TGGTTCTGCT	AATCTCCCAG GCTGGCAG	CA 120
TTGGAGACTT GCT	GACTTCT TTCATCC	CCC ACTCTTTCA	CCTGAAATTC CTTTCCTT	GG 180
TTTTGCTCTA AGI	CCTATGC TTCAGTC	AGG GGCCAACCAA	ATCTCACTGC CTCCTTTT	ra 240
TCATGAAGCC TTT	GATCACT GATAGTTO	CTT TTTATATCTT	GAAAAATCAC CCTTCCCA	300 gt
ACAGTTAATA TTT	PAGTATCT CTACTCAT	TCT TGGCACTTAC	TCACAGCTCC ATAATTCA	GT 360
CGTTTTCGTA CCT	CTTCATG GTGATGGG	GGA GCCCTTTGGA	GGTGGTGACT GTGCTTTA	ra 420
CTCCTCATGA TGC	CTTCACAT GTGGCAGG	GCG TGGAGTGCCC	GGAGGCGCC CTCCTGAT	rc 480
TGGGGCCTCC CAG			CCA GGC CCT GGG GGG Pro Gly Pro Gly Gly 10	530
	eu Leu Pro Ser Me		CTG CTG CTC CTG CTG Leu Leu Leu Leu 25	578
			TAC AAG GTG CCG GAG Tyr Lys Val Pro Glu 40	626
			GCA GCC GAC TAT GGT Ala Ala Asp Tyr Gly 60	674
			GTG GGT GCC CCG TAC Val Gly Ala Pro Tyr 75	722
Leu Arg Val As			ACC ACC GAG ACC TCC Thr Thr Glu Thr Ser 90	770
		lu Cys Gln Asn	CAG CTC CCT GGT GAT Gln Leu Pro Gly Asp 105	818
CCC TGC ATC CT Pro Cys Ile Le 110	GG GAG TTT GAG GT eu Glu Phe Glu Va 115	TA TCT ATC ACA	GAC CTC GTG CAG AAT Asp Leu Val Gln Asn 120	866
GCG AGC CCC CG Ala Ser Pro Ar 125	G CTG CTA GAG GG G Leu Leu Glu Gl 130	CC CAG ATA GAA Ly Gln Ile Glu 135	GTA CAA GAC ATC AAT Val Gln Asp Ile Asn 140	914
			ACT CTG GCC ATC CCT Thr Leu Ala Ile Pro 155	962
GAG AAC ACC AA Glu Asn Thr As 16	n Ile Gly Ser Le	TC TTC CCC ATC Eu Phe Pro Ile 165	CCG CTG GCT TCA GAC Pro Leu Ala Ser Asp 170	1010

CGT Arg	GAT Asp	GCT Ala 175	GGT Gly	CCC Pro	AAC Asn	GGT Gly	GTG Val 180	GCA Ala	TCC Ser	TAT Tyr	GAG Glu	CTG Leu 185	CAG Gln	GTG Val	GCA Ala	1058
		CAG Gln														1106
		GAG Glu														1154
		AGC Ser														1202
CTT	GAC Asp	ACC Thr	AAT Asn 240	GAC Asp	AAC Asn	GCC Ala	CCC Pro	AAG Lys 245	TTT Phe	GAG Glu	CGG Arg	CCC Pro	TCC Ser 250	TAT Tyr	GAG Glu	1250
GCC Ala	GAA Glu	CTA Leu 255	TCT Ser	GAG Glu	AAT Asn	AGC Ser	CCC Pro 260	ATA Ile	GGC Gly	CAC His	TCG Ser	GTC Val 265	ATC Ile	CAG Gln	GTG Val	1298
AAG Lys	GCC Ala 270	AAT Asn	GAC Asp	TCA Ser	GAC Asp	CAA Gln 275	GGT Gly	GCC Ala	AAT Asn	GCA Ala	GAA Glu 280	ATC Ile	GAA Glu	TAC Tyr	ACA Thr	1346
TTC Phe 285	CAC His	CAG Gln	GCG Ala	CCC Pro	GAA Glu 290	GTT Val	GTG Val	AGG Arg	CGT Arg	CTT Leu 295	CTT Leu	CGA Arg	CTG Leu	GAC Asp	AGG Arg 300	1394
AAC Asn	ACT Thr	GGA Gly	CTT Leu	ATC Ile 305	ACT Thr	GTT Val	CAG Gln	GGC Gly	CCG Pro 310	GTG Val	GAC Asp	CGT Arg	GAG Glu	GAC Asp 315	CTA Leu	1442
AGC Ser	ACC Thr	CTG Leu	CGC Arg 320	TTC Phe	TCA Ser	GTG Val	CTT Leu	GCT Ala 325	AAG Lys	GAC Asp	CGA Arg	GGC Gly	ACC Thr 330	AAC Asn	ccc Pro	1490
AAG Lys	AGT Ser	GCC Ala 335	CGT Arg	GCC Ala	CAG Gln	GTG Val	GTT Val 340	GTG Val	ACC Thr	GTG Val	AAG Lys	GAC Asp 345	ATG Met	AAT Asn	GAC Asp	1538
AAT Asn	GCC Ala 350	CCC Pro	ACC Thr	ATT Ile	GAG Glu	ATC Ile 355	CGG Arg	GGC Gly	ATA Ile	GGG Gly	CTA Leu 360	GTG Val	ACT Thr	CAT His	CAA Gln	1586
GAT Asp 365	GGG Gly	ATG Met	GCT Ala	AAC Asn	ATC Ile 370	TCA Ser	GAG Glu	GAT Asp	GTG Val	GCA Ala 375	GAG Glu	GAG Glu	ACA Thr	GCT Ala	GTG Val 380	1634
GCC Ala	CTG Leu	GTG Val	CAG Gln	GTG Val 385	TCT Ser	GAC Asp	CGA Arg	GAT Asp	GAG Glu 390	GGA Gly	GAG Glu	AAT Asn	GCA Ala	GCT Ala 395	GTC Val	1682
ACC Thr	TGT Cys	GTG Val	GTG Val 400	GCA Ala	GGT Gly	GAT Asp	GTG Val	CCC Pro 405	TTC Phe	CAG Gln	CTG Leu	CGC Arg	CAG Gln 410	GCC Ala	AGT Ser	1730

GAG Glu	ACA Thr	GGC Gly 415	Ser	GAC Asp	AGC Ser	AAG Lys	AAG Lys 420	AAG Lys	TAT Tyr	TTC Phe	CTG Leu	CAG Gln 425	ACT Thr	ACC Thr	ACC Thr	1778
CCG Pro	CTA Leu 430	Asp	TAC	GAG Glu	AAG Lys	GTC Val 435	AAA Lys	GAC Asp	TAC Tyr	ACC Thr	ATT Ile 440	GAG Glu	ATT Ile	GTG Val	GCT Ala	1826
GTG Val 445	GAC	TCT Ser	GGC Gly	AAC Asn	CCC Pro 450	CCA Pro	CTC	TCC Ser	AGC Ser	ACT Thr 455	AAC Asn	TCC Ser	CTC Leu	AAG Lys	GTG Val 460	1874
CAG Gln	GTG Val	GTG Val	GAC Asp	GTC Val 465	AAT Asn	GAC Asp	AAC Asn	GCA Ala	CCT Pro 470	GTC Val	TTC Phe	ACT Thr	CAG Gln	AGT Ser 475	GTC Val	1922
ACT Thr	GAG Glu	GTC Val	GCC Ala 480	TTC Phe	CCG Pro	GAA Glu	AAC Asn	AAC Asn 485	AAG Lys	CCT Pro	GGT Gly	GAA Glu	GTG Val 490	ATT Ile	GCT Ala	1970
GAG Glu	ATC Ile	ACT Thr 495	GCC Ala	AGT Ser	GAT Asp	GCT Ala	GAC Asp 500	TCT Ser	GGC Gly	TCT Ser	AAT Asn	GCT Ala 505	GAG Glu	CTG Leu	GTT Val	2018
TAC Tyr	TCT Ser 510	CTG Leu	GAG Glu	CCT Pro	GAG Glu	CCG Pro 515	GCT Ala	GCT Ala	AAG Lys	GGC Gly	CTC Leu 520	TTC Phe	ACC Thr	ATC Ile	TCA Ser	2066
CCC Pro 525	GAG Glu	ACT Thr	GGA Gly	GAG Glu	ATC Ile 530	CAG Gln	GTG Val	AAG Lys	ACA Thr	TCT Ser 535	CTG Leu	GAT Asp	CGG Arg	GAA Glu	CAG Gln 540	2114
cgg Arg	GAG Glu	AGC Ser	TAT Tyr	GAG Glu 545	TTG Leu	AAG Lys	GTG Val	GTG Val	GCA Ala 550	GCT Ala	GAC Asp	CGG Arg	GGC Gly	AGT Ser 555	CCT Pro	2162
AGC Ser	CTC Leu	CAG Gln	GGC Gly 560	ACA Thr	GCC Ala	ACT Thr	GTC Val	CTT Leu 565	GTC Val	AAT Asn	GTG Val	CTG Leu	GAC Asp 570	TGC Cys	AAT Asn	2210
GAC Asp	AAT Asn	GAC Asp 575	CCC Pro	AAA Lys	TTT Phe	ATG Met	CTG Leu 580	AGT Ser	GGC Gly	TAC Tyr	AAC Asn	TTC Phe 585	TCA Ser	GTG Val	ATG Met	2258
GAG Glu	AAC Asn 590	ATG Met	CCA Pro	GCA Ala	CTG Leu	AGT Ser 595	CCA Pro	GTG Val	GGC Gly	ATG Met	GTG Val 600	ACT Thr	GTC Val	ATT Ile	GAT Asp	2306
GGA Gly 605	GAC Asp	AAG Lys	GGG Gly	GAG Glu	AAT Asn 610	GCC Ala	CAG Gln	GTG Val	CAG Gln	CTC Leu 615	TCA Ser	GTG Val	GAG Glu	CAG Gln	GAC Asp 620	2354
AAC Asn	GGT Gly	GAC Asp	TTT Phe	GTT Val 625	ATC Ile	CAG Gln	AAT Asn	GGC Gly	ACA Thr 630	GGC Gly	ACC Thr	ATC Ile	CTA Leu	TCC Ser 635	AGC Ser	2402
CTG Leu	AGC Ser	TTT Phe	GAT Asp 640	CGA Arg	GAG Glu	CAA Gln	CAA Gln	AGC Ser 645	ACC Thr	TAC Tyr	ACC Thr	TTC Phe	CAG Gln 650	CTG Leu	AAG Lys	2450

GCA Ala	A GTG	GAI Asp 655	Gly	GGC Gly	GTC Val	CCA Pro	CCT Pro 660	Arg	TCA Ser	GCT Ala	TAC	GTT Val 665	GGT Gly	GTC Val	ACC	2498
ATC Ile	AAT Asn 670	Val	CTG Leu	GAC Asp	GAG Glu	AAT Asn 675	GAC Asp	AAC Asn	GCA Ala	CCC	TAT Tyr 680	Ile	ACT	GCC Ala	Pro	2546
TCT Ser 685	Asn	ACC Thr	TCT Ser	CAC	AAG Lys 690	Leu	CTG Leu	ACC Thr	CCC Pro	CAG Gln 695	ACA Thr	CGT Arg	CTT Leu	GGT Gly	GAG Glu 700	2594
ACG Thr	GTC Val	AGC Ser	CAG Gln	GTG Val 705	GCA Ala	GCC Ala	GAG Glu	GAC	TTT Phe 710	GAC Asp	TCT Ser	GGT Gly	GTC Val	AAT Asn 715	GCC Ala	2642
GAG Glu	CTG Leu	ATC Ile	TAC Tyr 720	AGC Ser	ATT	GCA Ala	GGT Gly	GGC Gly 725	AAC Asn	CCT Pro	TAT Tyr	GGA Gly	CTC Leu 730	TTC Phe	CAG Gln	2690
ATT Ile	GGG Gly	TCA Ser 735	CAT His	TCA Ser	GGT Gly	GCC Ala	ATC Ile 740	ACC Thr	CTG Leu	GAG Glu	AAG Lys	GAG Glu 745	ATT Ile	GAG Glu	CGG Arg	2738
CGC Arg	CAC His 750	CAT His	GGG Gly	CTA Leu	CAC His	CGC Arg 755	CTG Leu	GTG Val	GTG Val	AAG Lys	GTC Val 760	AGT Ser	GAC Asp	CGC Arg	GGC	2786
AAG Lys 765	CCC Pro	CCA Pro	CGC Arg	TAT Tyr	GGC Gly 770	ACA Thr	GCC Ala	TTG Leu	GTC Val	CAT His 775	CTT Leu	TAT Tyr	GTC Val	AAT Asn	GAG Glu 780	2834
ACT Thr	CTG Leu	GCC Ala	AAC Asn	CGC Arg 785	ACG Thr	CTG Leu	CTG Leu	GAG Glu	ACC Thr 790	CTC Leu	CTG Leu	GGC Gly	CAC His	AGC Ser 795	CTG Leu	2882
GAC Asp	ACG Thr	CCG Pro	CTG Leu 800	GAT Asp	ATT Ile	GAC Asp	ATT Ile	GCT Ala 805	GGG Gly	GAT Asp	CCA Pro	GAA Glu	TAT Tyr 810	GAG Glu	CGC Arg	2930
TCC Ser	AAG Lys	CAG Gln 815	CGT Arg	GGC Gly	AAC Asn	ATT Ile	CTC Leu 820	TTT Phe	GGT Gly	GTG Val	GTG Val	GCT Ala 825	GGT Gly	GTG Val	GTG Val	2978
GCC Ala	GTG Val 830	GCC Ala	TTG Leu	CTC Leu	ATC Ile	GCC Ala 835	CTG Leu	GCG Ala	GTT Val	CTT Leu	GTG Val 840	CGC Arg	TAC Tyr	TGC Cys	AGA Arg	3026
CAG Gln 845	CGG Arg	GAG Glu	GCC Ala	AAA Lys	AGT Ser 850	GGT Gly	TAC Tyr	CAG Gln	GCT Ala	GGT Gly 855	AAG Lys	AAG Lys	GAG Glu	ACC Thr	AAG Lys 860	3074
usp	Leu	TYE	ATA	865	гàв	Pro	Ser	Gly	AAG Lys 870	Ala	Ser	Lys	Gly	Asn 875	Lys	3122
AGC Ser	AAA Lys	GIŽ	AAG Lys 880	AAG Lys	AGC Ser	AAG Lys	Ser	CCA Pro 885	AAG Lys	CCC Pro	GTG Val	Lys	CCA Pro 890	GTG Val	GAG Glu	3170

GAC GAG GAT GAG GCC GGG CTG CAG AAG TCC CTC AAG TTC AAC CTG ATG Asp Glu Asp Glu Ala Gly Leu Gln Lys Ser Leu Lys Phe Asn Leu Met 895 900 905	3218
AGC GAT GCC CCT GGG GAC AGT CCC CGC ATC CAC CTG CCC CTC AAC TAC Ser Asp Ala Pro Gly Asp Ser Pro Arg Ile His Leu Pro Leu Asn Tyr 910 915 920	3266
CCA CCA GGC AGC CCT GAC CTG GGC CGC CAC TAT CGC TCT AAC TCC CCA Pro Pro Gly Ser Pro Asp Leu Gly Arg His Tyr Arg Ser Asn Ser Pro 925 930 940	3314
CTG CCT TCC ATC CAG CTG CAG CCC CAG TCA CCC TCA GCC TCC AAG AAG Leu Pro Ser Ile Gln Leu Gln Pro Gln Ser Pro Ser Ala Ser Lys Lys 945 950 955	3362
CAC CAG GTG GTA CAG GAC CTG CCA CCT GCA AAC ACA TTC GTG GGC ACC His Gln Val Val Gln Asp Leu Pro Pro Ala Asn Thr Phe Val Gly Thr 960 965 970	3410
GGG GAC ACC ACG TCC ACG GGC TCT GAG CAG TAC TCC GAC TAC AGC TAC Gly Asp Thr Thr Ser Thr Gly Ser Glu Gln Tyr Ser Asp Tyr Ser Tyr 980 985	3458
CGC ACC AAC CCC CCC AAA TAC CCC AGC AAG CAG GTA GGC CAG CCC TTT Arg Thr Asn Pro Pro Lys Tyr Pro Ser Lys Gln Val Gly Gln Pro Phe 990 995 1000	3506
CAG CTC AGC ACA CCC CAG CCC CTA CCC CAC CCC TAC CAC GGA GCC ATC Gln Leu Ser Thr Pro Gln Pro Leu Pro His Pro Tyr His Gly Ala Ile 1005 1010 1015 1020	3554
TGG ACC GAG GTG TGG GAG TGATGGAGCA GGTTTACTGT GCCTGCCCGT Trp Thr Glu Val Trp Glu 1025	3602
GTTGGGGGCC AGCCTGAGCC AGCAGTGGGA GGTGGGGCCT TAGTGCCTCA CCGGGCACAC	3662
GGATTAGGCT GAGTGAAGAT TAAGGGAGGG TGTGCTCTGT GGTCTCCTCC CTGCCCTCTC	3722
CCCACTGGGG AGAGACCTGT GATTTGCCAA GTCCCTGGAC CCTGGACCAG CTACTGGGCC	3782
TTATGGGTTG GGGGTAGG GCAGGTGAGC GTAAGTGGGG AGGGAAATGG GTAAGAAGTC	3842
TACTCCAAAC CTAGGTCTCT ATGTCAGACC AGACCTAGGT GCTTCTCTAG GAGGGAAACA	3902
GGGAGACCTG GGGTCCTGTG GATAACTGAG TGGGGAGTCT GCCAGGGGAG GGCACCTTCC	3962
CATTGTGCCT TCTGTGTGTA TTGTGCATTA ACCTCTTCCT CACCACTAGG CTTCTGGGGC	4022
TGGGTCCCAC ATGCCCTTGA CCCTGACAAT AAAGTTCTCT ATTTTTGGAA AAAAAAAAA	4082
AAAAAAAA AAAAAAAAA AA	4104

(2) INFORMATION FOR SEQ ID NO:95:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1026 amino acids
 - (B) TYPE: amino acid
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:95:

Met Glu Pro Leu Arg His Ser Pro Gly Pro Gly Gln Arg Leu Leu
1 5 10 15

Leu Pro Ser Met Leu Leu Ala Leu Leu Leu Leu Leu Ala Pro Ser Pro 20 25 30

Gly His Ala Thr Arg Val Val Tyr Lys Val Pro Glu Glu Gln Pro Pro 35 40 45

Asn Thr Leu Ile Gly Ser Leu Ala Ala Asp Tyr Gly Phe Pro Asp Val 50 55 60

Gly His Leu Tyr Lys Leu Glu Val Gly Ala Pro Tyr Leu Arg Val Asp 65 70 75 80

Gly Lys Thr Gly Asp Ile Phe Thr Thr Glu Thr Ser Ile Asp Arg Glu 85 90 95

Gly Leu Arg Glu Cys Gln Asn Gln Leu Pro Gly Asp Pro Cys Ile Leu 100 105 110

Glu Phe Glu Val Ser Ile Thr Asp Leu Val Gln Asn Ala Ser Pro Arg 115 120 125

Leu Leu Glu Gly Gln Ile Glu Val Gln Asp Ile Asn Asp Asn Thr Pro 130 135 140

Asn Phe Ala Ser Pro Val Ile Thr Leu Ala Ile Pro Glu Asn Thr Asn 145 150 155 160

Ile Gly Ser Leu Phe Pro Ile Pro Leu Ala Ser Asp Arg Asp Ala Gly 165 170 175

Pro Asn Gly Val Ala Ser Tyr Glu Leu Gln Val Ala Glu Asp Gln Glu 180 185 190

Glu Lys Gln Pro Gln Leu Ile Val Met Gly Asn Leu Asp Arg Glu Arg 195 200 205

Trp Asp Ser Tyr Asp Leu Thr Ile Lys Val Gln Asp Gly Gly Ser Pro 210 215 220

Pro Arg Ala Thr Ser Ala Leu Leu Arg Val Thr Val Leu Asp Thr Asn 225 230 235 240

Asp Asn Ala Pro Lys Phe Glu Arg Pro Ser Tyr Glu Ala Glu Leu Ser 245 250 255

Glu Asn Ser Pro Ile Gly His Ser Val Ile Gln Val Lys Ala Asn Asp 260 265 270

Ser Asp Gln Gly Ala Asn Ala Glu Ile Glu Tyr Thr Phe His Gln Ala 280 Pro Glu Val Val Arg Arg Leu Leu Arg Leu Asp Arg Asn Thr Gly Leu Ile Thr Val Gln Gly Pro Val Asp Arg Glu Asp Leu Ser Thr Leu Arg Phe Ser Val Leu Ala Lys Asp Arg Gly Thr Asn Pro Lys Ser Ala Arg Ala Gln Val Val Val Thr Val Lys Asp Met Asn Asp Asn Ala Pro Thr Ile Glu Ile Arg Gly Ile Gly Leu Val Thr His Gln Asp Gly Met Ala Asn Ile Ser Glu Asp Val Ala Glu Glu Thr Ala Val Ala Leu Val Gln Val Ser Asp Arg Asp Glu Gly Glu Asn Ala Ala Val Thr Cys Val Val Ala Gly Asp Val Pro Phe Gln Leu Arg Gln Ala Ser Glu Thr Gly Ser Asp Ser Lys Lys Lys Tyr Phe Leu Gln Thr Thr Thr Pro Leu Asp Tyr Glu Lys Val Lys Asp Tyr Thr Ile Glu Ile Val Ala Val Asp Ser Gly Asn Pro Pro Leu Ser Ser Thr Asn Ser Leu Lys Val Gln Val Val Asp Val Asn Asp Asn Ala Pro Val Phe Thr Gln Ser Val Thr Glu Val Ala Phe Pro Glu Asn Asn Lys Pro Gly Glu Val Ile Ala Glu Ile Thr Ala Ser Asp Ala Asp Ser Gly Ser Asn Ala Glu Leu Val Tyr Ser Leu Glu Pro Glu Pro Ala Ala Lys Gly Leu Phe Thr Ile Ser Pro Glu Thr Gly Glu Ile Gln Val Lys Thr Ser Leu Asp Arg Glu Gln Arg Glu Ser Tyr Glu Leu Lys Val Val Ala Ala Asp Arg Gly Ser Pro Ser Leu Gln Gly Thr Ala Thr Val Leu Val Asn Val Leu Asp Cys Asn Asp Asn Asp Pro 570 Lys Phe Met Leu Ser Gly Tyr Asn Phe Ser Val Met Glu Asn Met Pro

Ala Leu Ser Pro Val Gly Met Val Thr Val Ile Asp Gly Asp Lys Gly Glu Asn Ala Gln Val Gln Leu Ser Val Glu Gln Asp Asn Gly Asp Phe Val Ile Gln Asn Gly Thr Gly Thr Ile Leu Ser Ser Leu Ser Phe Asp Arg Glu Gln Gln Ser Thr Tyr Thr Phe Gln Leu Lys Ala Val Asp Gly Gly Val Pro Pro Arg Ser Ala Tyr Val Gly Val Thr Ile Asn Val Leu Asp Glu Asn Asp Asn Ala Pro Tyr Ile Thr Ala Pro Ser Asn Thr Ser His Lys Leu Leu Thr Pro Gln Thr Arg Leu Gly Glu Thr Val Ser Gln Val Ala Ala Glu Asp Phe Asp Ser Gly Val Asn Ala Glu Leu Ile Tyr
710 715 720 Ser Ile Ala Gly Gly Asn Pro Tyr Gly Leu Phe Gln Ile Gly Ser His 725 730 735 Ser Gly Ala Ile Thr Leu Glu Lys Glu Ile Glu Arg Arg His His Gly Leu His Arg Leu Val Val Lys Val Ser Asp Arg Gly Lys Pro Pro Arg 755 760 765 Tyr Gly Thr Ala Leu Val His Leu Tyr Val Asn Glu Thr Leu Ala Asn
770 780 Arg Thr Leu Leu Glu Thr Leu Leu Gly His Ser Leu Asp Thr Pro Leu 790 Asp Ile Asp Ile Ala Gly Asp Pro Glu Tyr Glu Arg Ser Lys Gln Arg Gly Asn Ile Leu Phe Gly Val Val Ala Gly Val Val Ala Val Ala Leu Leu Ile Ala Leu Ala Val Leu Val Arg Tyr Cys Arg Gln Arg Glu Ala Lys Ser Gly Tyr Gln Ala Gly Lys Lys Glu Thr Lys Asp Leu Tyr Ala 850 860 Pro Lys Pro Ser Gly Lys Ala Ser Lys Gly Asn Lys Ser Lys Gly Lys 865 Lys Ser Lys Ser Pro Lys Pro Val Lys Pro Val Glu Asp Glu Asp Glu 890 Ala Gly Leu Gln Lys Ser Leu Lys Phe Asn Leu Met Ser Asp Ala Pro

905

									- 19	•					
Gly	Asp	Ser 915	Pro	Arg	Ile	His	Leu 920	Pro	Leu	Asn	Tyr	Pro 925	Pro	Gly	Sei
Pro	Asp 930	Leu	Gly	Arg	His	Tyr 935	Arg	Ser	Asn	Ser	Pro 940	Leu	Pro	Ser	Ile
Gln 945	Leu	Gln	Pro	Gln	Ser 950	Pro	Ser	Ala	Ser	Lys 955	Lys	His	Gln	Val	Va]
Gln	Asp	Leu	Pro	Pro 965	Ala	Asn	Thr	Phe	Val 970	Gly	Thr	Gly	Asp	Thr 975	Thr
Ser	Thr	Gly	Ser 980	Glu	Gln	Tyr	Ser	Asp 985	Tyr	Ser	Tyr	Arg	Thr 990	Asn	Pro
Pro	Lys	Tyr 995	Pro	Ser	Lys	Gln	Val 1000	Gly	Gln	Pro	Phe	Gln 1005	Leu	Ser	Thr
Pro	Gln 1010	Pro	Leu	Pro	His	Pro 1015	Tyr	His	Gly	Ala	Ile 1020	Trp	Thr	Glu	Val
Trp 1025															
(2)	INFO	RMAT	ION	FOR	SEQ	ID N	0:96	:							

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 4705 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA
- (ix) FEATURE:

 - (A) NAME/KEY: CDS (B) LOCATION: 115..2827
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:96:

								-			•					
CGA	AAGC	CAT	GTCG	GACT	CG T	CGCC	CAGC	G CC	CAAG	CGCT	AAC	CCGC	TGA	AAGT	TTCTCA	60
GCG	AAAT	CTC .	AGGG.	ACGA'	TC T	GGAC	CCCG	C TG	AGAG	GAAC	TGC	TTTT	GAG	TGAG	ATG Met 1	117
GTC Val	CCA Pro	GAG Glu	GCC Ala 5	TGG Trp	AGG Arg	AGC Ser	GGA Gly	CTG Leu 10	GTA Val	AGC Ser	ACC Thr	GGG Gly	AGG Arg 15	GTA Val	GTG Val	165
GGA Gly	GTT Val	TTG Leu 20	CTT Leu	CTG Leu	CTT Leu	GGT Gly	GCC Ala 25	TTG Leu	AAC Asn	AAG Lys	GCT Ala	TCC Ser 30	ACG Thr	GTC Val	ATT Ile	213
CAC His	TAT Tyr 35	GAG Glu	ATC Ile	CCG Pro	GAG Glu	GAA Glu 40	AGA Arg	GAG Glu	AAG Lys	GGT Gly	TTC Phe 45	GCT Ala	GTG Val	GGC Gly	AAC Asn	261

GTG Val 50	Val	GCG Ala	AAC Asn	CTT Leu	GGT Gly 55	TTG Leu	GAT Asp	CTC Leu	GGT Gly	AGC Ser 60	CTC Leu	TCA Ser	GCC Ala	CGC Arg	AGG Arg 65	309
TTC Phe	CCG Pro	GTG Val	GTG Val	TCT Ser 70	GGA Gly	GCT Ala	AGC Ser	CGA Arg	AGA Arg 75	TTC Phe	TTT Phe	GAG Glu	GTG Val	AAC Asn 80	CGG Arg	357
GAG Glu	ACC Thr	GGA Gly	GAG Glu 85	ATG Met	TTT	GTG Val	AAC Asn	GAC Asp 90	CGT	CTG Leu	GAT Asp	CGA Arg	GAG Glu 95	GAG Glu	CTG Leu	405
TGT Cys	GGG Gly	ACA Thr 100	Leu	CCC Pro	TCT	TGC Cys	ACT Thr 105	GTA Val	ACT Thr	CTG Leu	GAG Glu	TTG Leu 110	GTA Val	GTG Val	GAG Glu	453
AAC Asn	CCG Pro 115	CTG Leu	GAG Glu	CTG Leu	TTC Phe	AGC Ser 120	GTG Val	GAA Glu	GTG Val	GTG Val	ATC Ile 125	CAG Gln	GAC Asp	ATC Ile	AAC Asn	501
GAC Asp 130	AAC Asn	AAT Asn	CCT Pro	GCT Ala	TTC Phe 135	CCT Pro	ACC Thr	CAG Gln	GAA Glu	ATG Met 140	AAA Lys	TTG Leu	GAG Glu	ATT Ile	AGC Ser 145	549
GAG Glu	GCC Ala	GTG Val	GCT Ala	CCG Pro 150	GGG Gly	ACG Thr	CGC Arg	TTT Phe	CCG Pro 155	CTC Leu	GAG Glu	AGC Ser	GCG Ala	CAC His 160	GAT Asp	597
CCC Pro	GAT Asp	CTG Leu	GGA Gly 165	AGC Ser	AAC Asn	TCT Ser	TTA Leu	CAA Gln 170	ACC Thr	TAT Tyr	GAG Glu	CTG Leu	AGC Ser 175	CGA Arg	AAT Asn	645
Glu	Tyr	Phe 180	Ala	Leu	Arg	GTG Val	Gln 185	Thr	Arg	Glu	Asp	Ser 190	Thr	Lys	Tyr	693
GCG Ala	GAG Glu 195	CTG Leu	GTG Val	TTG Leu	GAG Glu	CGC Arg 200	GCC Ala	CTG Leu	GAC Asp	CGA Arg	GAA Glu 205	CGG Arg	GAG Glu	CCT Pro	AGT Ser	741
CTC Leu 210	CAG Gln	TTA Leu	GTG Val	CTG Leu	ACG Thr 215	GCG Ala	TTG Leu	GAC Asp	GGA Gly	GGG Gly 220	ACC Thr	CCA Pro	GCT Ala	CTC Leu	TCC Ser 225	789
GCC Ala	AGC Ser	CTG Leu	CCT Pro	ATT Ile 230	CAC His	ATC Ile	AAG Lys	GTG Val	CTG Leu 235	GAC Asp	GCG Ala	AAT Asn	GAC Asp	AAT Asn 240	GCG Ala	837
CCT Pro	GTC Val	TTC Phe	AAC Asn 245	CAG Gln	TCC Ser	TTG Leu	TAC Tyr	CGG Arg 250	GCG Ala	CGC Arg	GTT Val	CCT Pro	GGA Gly 255	GGA Gly	TGC Cys	885
ACC Thr	TCC Ser	GGC Gly 260	ACG Thr	CGC Arg	GTG Val	GTA Val	CAA Gln 265	GTC Val	CTT Leu	GCA Ala	ACG Thr	GAT Asp 270	CTG Leu	GAT Asp	GAA Glu	933
GGC Gly	CCC Pro 275	AAC Asn	GGT Gly	GAA Glu	Ile	ATT Ile 280	TAC Tyr	TCC Ser	TTC Phe	GGC	AGC Ser 285	CAC His	AAC Asn	CGC Arg	GCC Ala	981

GG(G1 ₃ 29(Val	G CGC	G CAP	CTA Leu	TTC Phe 295	Ala	TTA Leu	GAC Asp	CTT Leu	GTA Val 300	Thr	GGG Gly	ATG Met	CTG Leu	ACA Thr 305		1029
ATO Ile	AAG Lys	GG1	CGG Arg	CTG Leu 310	Авр	TTC Phe	GAG Glu	GAC Asp	ACC Thr 315	AAA Lys	CTC	CAT His	GAG Glu	ATT Ile 320	TAC Tyr		1077
ATC Ile	CAG Gln	GCC Ala	AAA Lys 325	Asp	AAG Lys	GGC Gly	GCC Ala	AAT Asn 330	Pro	GAA Glu	GGA Gly	GCA Ala	CAT His 335	TGC Cys	AAA Lys		1125
GTG Val	TTG Leu	Val 340	Glu	GTT Val	GTG Val	GAT Asp	GTG Val 345	AAT Asn	GAC	AAC Asn	GCC Ala	CCG Pro 350	Glu	ATC Ile	ACA Thr		1173
GTC Val	Thr 355	Ser	GTG Val	TAC Tyr	AGC Ser	CCA Pro 360	GTA Val	CCC Pro	GAG Glu	GAT Asp	GCC Ala 365	TCT Ser	GGG Gly	ACT Thr	GTC Val		1221
370	Ala	Leu	CTC Leu	Ser	Val 375	Thr	Asp	Leu	Asp	Ala 380	Gly	Glu	Asn	Gly	Leu 385		1269
Val	Thr	Cys	GAA Glu	390	Pro	Pro	Gly	Leu	Pro 395	Phe	Ser	Leu	Thr	Ser 400	Ser		1317
Leu	rys	Asn	TAC Tyr 405	Phe	Thr	Leu	Lys	Thr 410	Ser	Ala	Asp	Leu	Asp 415	Arg	Glu		1365
Inr	Vai	420	GAA Glu	Tyr	Asn	Leu	Ser 425	Ile	Thr	Ala	Arg	Asp 430	Ala	Gly	Thr	:	1413
PIO	435	rea	TCA Ser	Ala	Leu	440	Ile	Val	Arg	Val	Gln 445	Val	Ser	Asp	Ile	:	1461
450	rap	ABII	CCT Pro	PIO	455	ser	ser	Gin	Ser	Ser 460	Tyr	Asp	Val	Tyr	Ile 465	2	1509
GIU	GIU	ABII	AAC Asn	470	Pro	GIĀ	Ala	Pro	11e 475	Leu	Asn	Leu	Ser	Val 480	Trp	1	1557
rsp	PIO	Asp	GCC Ala 485	Pro	GIN	Asn	Ala	Arg 490	Leu	Ser	Phe	Phe	Leu 495	Leu	Glu	1	1605
GIII	GLY	500	GAA Glu	THE	GIĀ	Leu	Val 505	GIÀ	Arg	Tyr	Phe	Thr 510	Ile	Asn	Arg	1	1653
GAC Asp	AAT Asn 515	GGC Gly	ATA Ile	GTG Val	ser	TCC Ser :	TTA Leu	GTG Val	CCC Pro	Leu	GAC Asp 525	TAT Tyr	GAG Glu	GAT Asp	CGG Arg	1	1701

CGG Arg 530	Glu	TTI Phe	GAA Glu	TTA Leu	ACA Thr 535	GCT Ala	CAT His	ATC Ile	AGC Ser	GAT Asp 540	GGG Gly	GGC	ACC Thr	CCG Pro	GTC Val 545	1749
CTA Leu	GCC Ala	ACC	AAC Asn	Ile 550	Ser	GTG Val	AAC Asn	ATA Ile	TTT Phe 555	GTC Val	ACT Thr	GAT Asp	CGC Arg	AAT Asn 560	GAC Asp	1797
AAT Asn	GCC	CCC	CAG Gln 565	GTC Val	CTA Leu	TAT Tyr	CCT Pro	CGG Arg 570	CCA Pro	GGT Gly	GGG Gly	AGC Ser	TCG Ser 575	GTG Val	GAG Glu	1845
ATG Met	CTG Leu	CCT Pro 580	Arg	GGT	ACC Thr	TCA Ser	GCT Ala 585	GGC Gly	CAC His	CTA Leu	GTG Val	TCA Ser 590	CGG Arg	GTG Val	GTA Val	1893
GGC Gly	TGG Trp 595	GAC Asp	GCG Ala	GAT Asp	GCA Ala	GGG Gly 600	CAC His	AAT Asn	GCC Ala	TGG Trp	CTC Leu 605	TCC Ser	TAC Tyr	AGT Ser	CTC Leu	1941
TTT Phe 610	GGA Gly	TCC Ser	CCT Pro	AAC Asn	CAG Gln 615	AGC Ser	CTT Leu	TTT Phe	GCC Ala	ATA Ile 620	GGG Gly	CTG Leu	CAC His	ACT Thr	GGT Gly 625	1989
CAA Gln	ATC Ile	AGT Ser	ACT Thr	GCC Ala 630	CGT Arg	CCA Pro	GTC Val	CAA Gln	GAC Asp 635	ACA Thr	GAT Asp	TCA Ser	CCC Pro	AGG Arg 640	CAG Gln	2037
ACT Thr	CTC Leu	ACT Thr	GTC Val 645	TTG Leu	ATC Ile	AAA Lys	GAC Asp	AAT Asn 650	GGG Gly	GAG Glu	CCT Pro	TCG Ser	CTC Leu 655	TCC Ser	ACC Thr	2085
ACT Thr	GCT Ala	ACC Thr 660	CTC Leu	ACT Thr	GTG Val	TCA Ser	GTA Val 665	ACC Thr	GAG Glu	GAC Asp	TCT Ser	CCT Pro 670	GAA Glu	GCC Ala	CGA Arg	2133
GCC Ala	GAG Glu 675	TTC Phe	CCC Pro	TCT Ser	GGC Gly	TCT Ser 680	GCC Ala	CCC Pro	CGG Arg	GAG Glu	CAG Gln 685	AAA Lys	AAA Lys	AAT Asn	CTC Leu	2181
ACC Thr 690	TTT Phe	TAT Tyr	CTA Leu	CTT Leu	CTT Leu 695	TCT Ser	CTA Leu	ATC Ile	CTG Leu	GTT Val 700	TCT Ser	GTG Val	GGC Gly	TTC Phe	GTG Val 705	2229
GTC Val	ACA Thr	GTG Val	TTC Phe	GGA Gly 710	GTA Val	ATC Ile	ATA Ile	TTC Phe	AAA Lys 715	GTT Val	TAC Tyr	AAG Lys	TGG Trp	AAG Lys 720	CAG Gln	2277
TCT Ser	AGA Arg	GAC Asp	CTA Leu 725	TAC Tyr	CGA Arg	GCC Ala	CCG Pro	GTG Val 730	AGC Ser	TCA Ser	CTG Leu	TAC Tyr	CGA Arg 735	ACA Thr	CCA Pro	2325
GGG Gly	CCC Pro	TCC Ser 740	TTG Leu	CAC His	GCG Ala	GAC Asp	GCC Ala 745	GTG Val	CGG Arg	GGA Gly	GGC Gly	CTG Leu 750	ATG Met	TCG Ser	CCG Pro	2373
CAC His	CTT Leu 755	TAC Tyr	CAT His	CAG Gln	Val	TAT Tyr 760	CTC Leu	ACC Thr	ACG Thr	Asp	TCC Ser 765	CGC Arg	CGC Arg	AGC Ser	GAC Asp	2421

CCG CTG CTG AAG AAA CCT GGT GCA GCC AGT CCA CTG GCC AGC CGC CAG Pro Leu Leu Lys Lys Pro Gly Ala Ala Ser Pro Leu Ala Ser Arg Gln 770 785	2469
AAC ACG CTG CGG AGC TGT GAT CCG GTG TTC TAT AGG CAG GTG TTG GGT Asn Thr Leu Arg Ser Cys Asp Pro Val Phe Tyr Arg Gln Val Leu Gly 790 795 800	2517
GCA GAG AGC GCC CCT CCC GGA CAG CAA GCC CCG CCC AAC ACG GAC TGG Ala Glu Ser Ala Pro Pro Gly Gln Gln Ala Pro Pro Asn Thr Asp Trp 805 810 815	2565
CGT TTC TCT CAG GCC CAG AGA CCC GGC ACC AGC GGC TCC CAA AAT GGC Arg Phe Ser Gln Ala Gln Arg Pro Gly Thr Ser Gly Ser Gln Asn Gly 820 825	2613
GAT GAC ACC GGC ACC TGG CCC AAC AAC CAG TTT GAC ACA GAG ATG CTG Asp Asp Thr Gly Thr Trp Pro Asn Asn Gln Phe Asp Thr Glu Met Leu 835	2661
CAA GCC ATG ATC TTG GCG TCC GCC AGT GAA GCT GCT GAT GGG AGC TCC Gln Ala Met Ile Leu Ala Ser Ala Ser Glu Ala Ala Asp Gly Ser Ser 850 865	2709
ACC CTG GGA GGG GGT GCC GGC ACC ATG GGA TTG AGC GCC CGC TAC GGA Thr Leu Gly Gly Ala Gly Thr Met Gly Leu Ser Ala Arg Tyr Gly 870 875 880	2757
CCC CAG TTC ACC CTG CAG CAC GTG CCC GAC TAC CGC CAG AAT GTC TAC Pro Gln Phe Thr Leu Gln His Val Pro Asp Tyr Arg Gln Asn Val Tyr 885 890 895	2805
ATC CCA GGC AGC AAT GCA CAC T GACCAACGCA GCTGGCAAGC GGATGGCAAG Ile Pro Gly Ser Asn Ala His 900	2857
GCCCAGCAGG TGGCAATGGC AACAAGAAGA AGTCGGCAAG AAGGAGAAGA AGTAACATGG	2917
AGGCCAGGCC AAGAGCCACA GGGCAGCCTC TCCCCGAACC AGCCCAGCTT CTCCTTACCT	2977
GCACCCAGGC CTCAGAGTTT CAGGGCTAAC CCCCAGAATA CTGGTAGGGG CCAAGGCATC	3037
TCCCTTGGAA ACAGAAACAA GTGCCATCAC ACCATCCCTT CCCCAGGTGT AATATCCAAA	3097
GCAGTTCCGC TGGGAACCCC ATCCAATCAG TGGCTGTACC CATTTGGGTA GTGGGGTTCA	3157
TGTAGACACC AAGAACCATT TGCCACACCC CGTTTAGTTA CAGCTGAACC CTCCATCTTC	3217
CAAATCAATC AGGCCCATCC ATCCCATGCC TCCCTCCTCC CCACCCCACT CCAACAGTTC	3277
CTCTTTCCCG AGTAAGGTGG TTGGGGTGTT GAAGTACCAA GTAACCTACA AGCCTCCTAG	3337
TTCTGAAAAG TTGGAAGGGC ATCATGACCT CTTGGCCTCT CCTTTGATTC TCAATCTTCC	3397
CCCAAAGCAT GGTTTGGTGC CAGCCCCTTC ACCTCCTTCC AGAGCCCAAG ATCAATGCTC	3457
AAGTTTTGGA GGACATGATC ACCATCCCCA TGGTACTGAT GCTTGCTGGA TTTAGGGAGG	3517
GCATTTTGCT ACCAAGCCTC TTCCCAACGC CCTGGGACCA GTCTTCTGTT TTGTTTTCA	3577
TTGTTTGAGC TTTCCACTGC ATGCCTTGAC TTCCCCCACC TCCTCCTCAA ACAAGAGACT	3637

	CCACTGCATG	TTCCAAGACA	. GTATGGGGTG	GTAAGATAAG	GAAGGGAAGT	GTGTGGATGT	3697
	GGATGGTGGG	GGCATGGACA	AAGCTTGACA	CATCAAGTTA	TCAAGGCCTT	GGAGGAGGCT	3757
	CTGTATGTCC	TCAGGGGACT	GACAACATCC	TCCAGATTCC	AGCCATAAAC	CAATAACTAG	3817
	GCTGGACCCT	TCCCACTACA	TAATAGGGCT	CAGCCAGGCA	GCCAGCTTTG	GGCTGAGCTA	3877
	ACAGGACCAA	TGGATTAACT	GGCATTTCAG	TCCAAGGAAG	CTCGAAGCAG	GTTTAGGACC	3937
٠	AGGTCCCCTT	GAGAGGTCAG	AGGGGCCTCT	GTGGGTGCTG	GGTACTCCAG	AGGTGCCACT	3997
(GGTGGAAGGG	TCAGCGGAGC	CCCAGCAGGA	AGGGTGGGCC	AGCCAGGCCA	TTCTTAGTCC	4057
(CTGGGTTGGG	GAGGCAGGGA	GCTAGGGCAG	GGACCAAATG	AACAGAAAGT	CTCAGCCCAG	4117
(GATGGGGCTT	CTTCAACAGG	CCCCTGCCCT	CCTGAAGCCT	CAGTCCTTCA	CCTTGCCAGG	4177
•	IGCCGTTTCT	CTTCCGTGAA	GGCCACTGCC	CAGGTCCCCA	GTGCGCCCCC	TAGTGGCCAT	4237
2	AGCCTGGTTA	AAGTTCCCCA	GTGCCTCCTT	GTGATAGACC	TTCTTCTCCC	ACCCCTTCT	4297
(SCCCCTGGGT	CCCCGGCCAT	CCAGCGGGGC	TGCCAGAGAA	CCCCAGACCT	GCCCTTACAG	4357
7	PAGTGTAGCG	CCCCCTCCCT	CTTTCGGCTG	GTGTAGAATA	GCCAGTAGTG	TAGTGCGGTG	4417
7	GCTTTTACG	TGATGGCGGG	TGGGCAGCGG	GCGGCGCGT	CCGCGCAGCC	GTCTGTCCTT	4477
C	SATCTGCCCG	CGGCGGCCCG	TGTTGTGTTT	TGTGCTGTGT	CCAGCGCTAA	GGCGACCCC	4537
7	CCCCCGTAC	TGACTTCTCC	TATAAGCGCT	TCTCTTCGCA	TAGTCACGTA	GCTCCCACCC	4597
C	ACCCTCTTC	CTGTGTCTCA	CGCAAGTTTT	ATACTCTAAT	ATTTATATGG	CTTTTTTCT	4657
1	CGACAAAAA	AATAATAAA	CGTTTCTTCT	GAAAAAAAA	AAAAAAA		4705

(2) INFORMATION FOR SEQ ID NO:97:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 904 amino acids

 - (B) TYPE: amino acid(D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: protein
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:97:
- Met Val Pro Glu Ala Trp Arg Ser Gly Leu Val Ser Thr Gly Arg Val
- Val Gly Val Leu Leu Leu Gly Ala Leu Asn Lys Ala Ser Thr Val
- Ile His Tyr Glu Ile Pro Glu Glu Arg Glu Lys Gly Phe Ala Val Gly
- Asn Val Val Ala Asn Leu Gly Leu Asp Leu Gly Ser Leu Ser Ala Arg 50 55 60

Arg Phe Pro Val Val Ser Gly Ala Ser Arg Arg Phe Phe Glu Val Asn Arg Glu Thr Gly Glu Met Phe Val Asn Asp Arg Leu Asp Arg Glu Glu Leu Cys Gly Thr Leu Pro Ser Cys Thr Val Thr Leu Glu Leu Val Val Glu Asn Pro Leu Glu Leu Phe Ser Val Glu Val Val Ile Gln Asp Ile Asn Asp Asn Asn Pro Ala Phe Pro Thr Gln Glu Met Lys Leu Glu Ile Ser Glu Ala Val Ala Pro Gly Thr Arg Phe Pro Leu Glu Ser Ala His Asp Pro Asp Leu Gly Ser Asn Ser Leu Gln Thr Tyr Glu Leu Ser Arg Asn Glu Tyr Phe Ala Leu Arg Val Gln Thr Arg Glu Asp Ser Thr Lys Tyr Ala Glu Leu Val Leu Glu Arg Ala Leu Asp Arg Glu Arg Glu Pro Ser Leu Gln Leu Val Leu Thr Ala Leu Asp Gly Gly Thr Pro Ala Leu Ser Ala Ser Leu Pro Ile His Ile Lys Val Leu Asp Ala Asn Asp Asn 230 Ala Pro Val Phe Asn Gln Ser Leu Tyr Arg Ala Arg Val Pro Gly Gly 245 250 255 Cys Thr Ser Gly Thr Arg Val Val Gln Val Leu Ala Thr Asp Leu Asp Glu Gly Pro Asn Gly Glu Ile Ile Tyr Ser Phe Gly Ser His Asn Arg Ala Gly Val Arg Gln Leu Phe Ala Leu Asp Leu Val Thr Gly Met Leu Thr Ile Lys Gly Arg Leu Asp Phe Glu Asp Thr Lys Leu His Glu Ile Tyr Ile Gln Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys Lys Val Leu Val Glu Val Val Asp Val Asp Asp Asp Asp Ala Pro Glu Ile Thr Val Thr Ser Val Tyr Ser Pro Val Pro Glu Asp Ala Ser Gly Thr 360 Val Ile Ala Leu Leu Ser Val Thr Asp Leu Asp Ala Gly Glu Asn Gly

375

Leu Val Thr Cys Glu Val Pro Pro Gly Leu Pro Phe Ser Leu Thr Ser 390 Ser Leu Lys Asn Tyr Phe Thr Leu Lys Thr Ser Ala Asp Leu Asp Arg Glu Thr Val Pro Glu Tyr Asn Leu Ser Ile Thr Ala Arg Asp Ala Gly Thr Pro Ser Leu Ser Ala Leu Thr Ile Val Arg Val Gln Val Ser Asp Ile Asn Asp Asn Pro Pro Gln Ser Ser Gln Ser Ser Tyr Asp Val Tyr 455 Ile Glu Glu Asn Asn Leu Pro Gly Ala Pro Ile Leu Asn Leu Ser Val Trp Asp Pro Asp Ala Pro Gln Asn Ala Arg Leu Ser Phe Phe Leu Leu Glu Gln Gly Ala Glu Thr Gly Leu Val Gly Arg Tyr Phe Thr Ile Asn Arg Asp Asn Gly Ile Val Ser Ser Leu Val Pro Leu Asp Tyr Glu Asp Arg Arg Glu Phe Glu Leu Thr Ala His Ile Ser Asp Gly Gly Thr Pro Val Leu Ala Thr Asn Ile Ser Val Asn Ile Phe Val Thr Asp Arg Asn 550 Asp Asn Ala Pro Gln Val Leu Tyr Pro Arg Pro Gly Gly Ser Ser Val Glu Met Leu Pro Arg Gly Thr Ser Ala Gly His Leu Val Ser Arg Val Val Gly Trp Asp Ala Asp Ala Gly His Asn Ala Trp Leu Ser Tyr Ser Leu Phe Gly Ser Pro Asn Gln Ser Leu Phe Ala Ile Gly Leu His Thr Gly Gln Ile Ser Thr Ala Arg Pro Val Gln Asp Thr Asp Ser Pro Arg 630 Gln Thr Leu Thr Val Leu Ile Lys Asp Asn Gly Glu Pro Ser Leu Ser 645 Thr Thr Ala Thr Leu Thr Val Ser Val Thr Glu Asp Ser Pro Glu Ala Arg Ala Glu Phe Pro Ser Gly Ser Ala Pro Arg Glu Gln Lys Lys Asn 680 Leu Thr Phe Tyr Leu Leu Leu Ser Leu Ile Leu Val Ser Val Gly Phe

Val Val Thr Val Phe Gly Val Ile Ile Phe Lys Val Tyr Lys Trp Lys

Gln Ser Arg Asp Leu Tyr Arg Ala Pro Val Ser Ser Leu Tyr Arg Thr

Pro Gly Pro Ser Leu His Ala Asp Ala Val Arg Gly Gly Leu Met Ser

Pro His Leu Tyr His Gln Val Tyr Leu Thr Thr Asp Ser Arg Arg Ser 755 760 765

Asp Pro Leu Leu Lys Lys Pro Gly Ala Ala Ser Pro Leu Ala Ser Arg

Gln Asn Thr Leu Arg Ser Cys Asp Pro Val Phe Tyr Arg Gln Val Leu 785

Gly Ala Glu Ser Ala Pro Pro Gly Gln Gln Ala Pro Pro Asn Thr Asp

Trp Arg Phe Ser Gln Ala Gln Arg Pro Gly Thr Ser Gly Ser Gln Asn

Gly Asp Asp Thr Gly Thr Trp Pro Asn Asn Gln Phe Asp Thr Glu Met 840

Leu Gln Ala Met Ile Leu Ala Ser Ala Ser Glu Ala Ala Asp Gly Ser

Ser Thr Leu Gly Gly Gly Ala Gly Thr Met Gly Leu Ser Ala Arg Tyr 865 870 875 880

Gly Pro Gln Phe Thr Leu Gln His Val Pro Asp Tyr Arg Gln Asn Val 890

Tyr Ile Pro Gly Ser Asn Ala His 900

(2) INFORMATION FOR SEQ ID NO:98:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 441 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:98:

Asp Trp Val Ile Pro Pro Ile Asn Leu Pro Glu Asn Ser Arg Gly Pro

Phe Pro Gln Glu Leu Val Arg Ile Arg Ser Asp Arg Asp Lys Asn Leu 20 25 30

Ser Leu Arg Tyr Thr Val Thr Gly Pro Gly Ala Asp Gln Pro Pro Thr 35 40 45

Gly Ile Phe Ile Ile Asn Pro Ile Ser Gly Gln Leu Ser Val Thr Lys
50 55 60

Pro Leu Asp Arg Glu Gln Ile Ala Arg Phe His Leu Arg Ala His Ala 65 70 75 80

Val Asp Ile Asn Gly Asn Gln Val Glu Asn Pro Ile Asp Ile Val Ile 85 90 95

Asn Val Ile Asp Met Asn Asp Asn Arg Pro Glu Phe Thr Ala Met Thr 100 105 110

Phe Tyr Gly Glu Val Pro Glu Asn Arg Val Asp Ile Ile Val Ala Asn 115 120 125

Leu Thr Val Thr Asp Lys Asp Gln Pro His Thr Pro Ala Trp Asn Ala 130 135 140

Val Thr Arg Ile Ser Gly Gly Asp Pro Thr Gly Arg Phe Ala Ile Gln 145 150 155 160

Thr Asp Pro Asn Ser Asn Asp Gly Leu Val Thr Val Val Lys Pro Ile 165 170 175

Asp Phe Glu Thr Asn Arg Met Phe Val Leu Thr Val Ala Ala Glu Asn 180 185 190

Gln Val Pro Leu Ala Lys Gly Ile Gln His Pro Pro Gln Ser Thr Ala 195 200 205

Thr Val Ser Val Thr Val Ile Asp Val Asn Glu Asn Pro Tyr Phe Ala 210 215 220

Pro Asn Pro Lys Ile Ile Arg Gln Glu Glu Gly Leu His Ala Gly Thr 230 235 240

Met Leu Thr Thr Phe Thr Ala Gly Asp Pro Asp Arg Tyr Met Gln Gln 245 250 255

Asn Ile Arg Tyr Thr Lys Leu Ser Asp Pro Ala Asn Trp Leu Lys Ile 260 265 270

Asp Pro Val Asn Gly Gln Ile Thr Thr Ile Ala Val Leu Asp Arg Glu 275

Ser Pro Asn Val Lys Asn Asn Ile Tyr Asn Ala Thr Phe Leu Ala Ser 290 295 300

Asp Asn Gly Ile Pro Pro Met Ser Gly Thr Gly Thr Leu Gln Ile Tyr 305 310 315 320

Leu Leu Asp Ile Asn Asp Asn Ala Pro Gln Val Leu Pro Gln Glu Ala 325 330 335

Glu Thr Cys Glu Thr Pro Asp Pro Asn Ser Ile Asn Ile Thr Thr Ala 340 345 350

Leu Asp Tyr Asp Ile Asp Pro Asn Ala Gly Pro Phe Ala Tyr Asp Leu

Pro Leu Ser Pro Val Thr Ile Lys Arg Asn Trp Thr Ile Thr Arg Leu

Asn Gly Asp Phe Ala Gln Leu Asn Leu Lys Ile Lys Phe Leu Glu Ala

Gly Ile Tyr Glu Val Pro Ile Ile Ile Thr Asp Ser Gly Asn Pro Pro

Lys Ser Asn Lys Ser Ile Leu Arg Val Arg Val Cys Gln Cys Asp Phe

Asn Gly Asp Cys Thr Asp Val Asp Arg

(2) INFORMATION FOR SEQ ID NO:99:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 105 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: protein
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:99:

Glu Asp Thr Val Tyr Ser Phe Asp Ile Pro Glu Asn Ala Gln Arg Gly

Tyr Gln Val Gly Gln Ile Val Ala Arg Asp Ala Asp Leu Gly Gln Asn

Ala Gln Leu Ser Tyr Gly Val Val Ser Asp Trp Ala Asn Asp Val Phe

Ser Leu Asn Pro Gln Thr Gly Met Leu Thr Leu Thr Ala Arg Leu Asp

Tyr Glu Glu Val Gln His Tyr Ile Leu Ile Val Gln Ala Gln Asp Asn 65 70 75 80

Gly Gln Pro Ser Leu Ser Thr Thr Ile Thr Val Tyr Cys Asn Val Leu

Asp Leu Asn Asp Asn Ala Pro Ile Phe 100

(2) INFORMATION FOR SEQ ID NO:100:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 7 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single (D) TOPOLOGY: linear

(ii)	MOLECULE	TYPE:	protein
------	----------	-------	---------

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:100:

Asp Xaa Asp Xaa Gly Xaa Asn

- (2) INFORMATION FOR SEQ ID NO:101:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 7 amino acids
 (B) TYPE: amino acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: protein
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:101:

Ala Xaa Asp Xaa Gly Xaa Pro

- (2) INFORMATION FOR SEQ ID NO:102:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 4650 base pairs

 - (B) TYPE: nucleic acid (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA
 - (ix) FEATURE:

 - (A) NAME/KEY: CDS (B) LOCATION: 495..4103
 - (xi) SEQUENCE DESCRIPTION: SEQ ID NO:102:

CCTCTATTCG	ACATTCTCTT	TGGATTGTTT	TGCTATAACT	TGAAATTTGG	GATGTCACAA	60
ACGAAACTGT	CATCTGTTTC	CGCCAAACTG	TGGTTCTGCT	AATCTCCCAG	GCTGGCAGCA	120
TTGGAGACTT	GCTGACTTCT	TTCATCCCCC	ACTCTTTTCA	CCTGAAATTC	CTTTCCTTGG	180
TTTTGCTCTA	AGTCCTATGC	TTCAGTCAGG	GGCCAACCAA	ATCTCACTGC	CTCCTTTTTA	240
TCATGAAGCC	TTTGATCACT	GATAGTTCTT	TTTATATCTT	GAAAAATCAC	CCTTCCCAGT	300
ACAGTTAATA	TTTAGTATCT	CTACTCATCT	TGGCACTTAC	TCACAGCTCC	ATAATTCAGT	360
CGTTTTCGTA	CCTCTTCATG	GTGATGGGGA	GCCCTTTGGA	GGTGGTGACT	GTGCTTTATA	420
CTCCTCATGA	TGCTTCACAT	GTGGCAGGCG	TGGAGTGCCC	GGAGGCGGCC	CTCCTGATTC	480

TGGGGCCTCC CAGG	ATG GAG Met Glu 1							530
CAA CGG CTA CTG Gln Arg Leu Leu 15								578
GCT CCA TCC CCA Ala Pro Ser Pro 30				l Val T				626
GAA CAG CCA CCC Glu Gln Pro Pro 45								674
TTT CCA GAT GTG Phe Pro Asp Val				ı Glu V				722
CTT CGC GTG GAT Leu Arg Val Asp 80								770
ATC GAC CGT GAG Ile Asp Arg Glu 95								818
CCC TGC ATC CTG Pro Cys Ile Leu 110				Thr A				866
GCG AGC CCC CGG Ala Ser Pro Arg 125	CTG CTA Leu Leu 130	GAG GGC Glu Gly	CAG ATI	GAA G Glu V 135	TA CAA al Gln	GAC ATC Asp Ile	AAT Asn 140	914
GAC AAC ACA CCC Asp Asn Thr Pro	AAC TTC Asn Phe 145	GCC TCA Ala Ser	CCA GTO Pro Val	Ile T	CT CTG	GCC ATC Ala Ile 155	CCT Pro	962
GAG AAC ACC AAC Glu Asn Thr Asn 160	ATC GGC Ile Gly	TCA CTC Ser Leu	TTC CCC Phe Pro 165	ATC CO	ro Leu	GCT TCA Ala Ser 170	GAC Asp	1010
CGT GAT GCT GGT Arg Asp Ala Gly 175	CCC AAC Pro Asn	GGT GTG Gly Val 180	GCA TCC Ala Ser	TAT G	AG CTG lu Leu 185	CAG GTG Gln Val	GCA Ala	1058
GAG GAC CAG GAG Glu Asp Gln Glu 190	GAG AAG Glu Lys	CAA CCA Gln Pro 195	CAG CTC	Ile Va	TG ATG (al Met (GGC AAC Gly Asn	CTG Leu	1106
GAC CGT GAG CGC Asp Arg Glu Arg 205	TGG GAC Trp Asp 210	TCC TAT Ser Tyr	GAC CTO Asp Lev	Thr II	TC AAG	GTG CAG Val Gln	GAT Asp 220	1154
GGC GGC AGC CCC Gly Gly Ser Pro	CCA CGC Pro Arg 225	GCC ACG Ala Thr	AGT GCC Ser Ala 230	Leu Le	TG CGT (GTC ACC Val Thr 235	GTG Val	1202

CTI Leu	Asp Asp	ACC Thr	AAT Asn 240	Asp	AAC Asn	GCC Ala	CCC Pro	AAG Lys 245	Phe	GAG Glu	CGG Arg	CCC Pro	TCC Ser 250	Tyr	GAG Glu	1250
GCC Ala	GAA Glu	CTA Leu 255	Ser	GAG Glu	AAT Asn	AGC Ser	CCC Pro 260	ATA Ile	GGC	CAC His	TCG Ser	GTC Val 265	ATC Ile	CAG Gln	GTG Val	1298
AAG Lys	GCC Ala 270	Asn	GAC Asp	TCA Ser	GAC Asp	CAA Gln 275	GGT Gly	GCC Ala	AAT Asn	GCA Ala	GAA Glu 280	ATC Ile	GAA Glu	TAC Tyr	ACA Thr	1346
TTC Phe 285	His	CAG Gln	GCG Ala	CCC Pro	GAA Glu 290	GTT Val	GTG Val	AGG Arg	CGT	CTT Leu 295	CTT	CGA Arg	CTG Leu	GAC Asp	AGG Arg 300	1394
AAC Asn	ACT Thr	GGA Gly	CTT Leu	ATC Ile 305	ACT Thr	GTT Val	CAG Gln	GGC Gly	CCG Pro 310	GTG Val	GAC Asp	CGT Arg	GAG Glu	GAC Asp 315	CTA Leu	1442
AGC Ser	ACC Thr	CTG Leu	CGC Arg 320	TTC Phe	TCA Ser	GTG Val	CTT Leu	GCT Ala 325	AAG Lys	GAC Asp	CGA Arg	GGC Gly	ACC Thr 330	AAC Asn	CCC Pro	1490
Lys	Ser	Ala 335	Arg	Ala	Gln	Val	Val 340	Val	ACC Thr	Val	Lys	Asp 345	Met	Asn	Asp	1538
AAT Asn	GCC Ala 350	CCC Pro	ACC Thr	ATT	GAG Glu	ATC Ile 355	CGG Arg	GGC Gly	ATA Ile	GGG Gly	CTA Leu 360	GTG Val	ACT Thr	CAT His	CAA Gln	1586
365	Gly	Met	Ala	Asn	11e 370	Ser	Glu	Asp	GTG Val	Ala 375	Glu	Glu	Thr	Ala	Val 380	1634
Ala	Leu	Val	Gln	Val 385	Ser	Asp	Arg	yeb	GAG Glu 390	Gly	Glu	Asn	Ala	Ala 395	Val	1682
Thr	Cys	Val	Val 400	Ala	Gly	Asp	Val	Pro 405	TTC Phe	Gln	Leu	Arg	Gln 410	Ala	Ser	1730
GIu	Thr	Gly 415	Ser	Asp	Ser	Lys	Lys 420	Lys	TAT Tyr	Phe	Leu	Gln 425	Thr	Thr	Thr	1778
Pro	430	Asp	Tyr	Glu	Lys	Val 435	Lys	Asp	TAC Tyr	Thr	11e 440	Glu	Ile	Val	Ala	1826
Val 445	Asp	Ser	Gly	Asn	Pro 450	Pro	Leu	Ser		Thr 455	Asn	Ser	Leu	Lys	Val 460	1874
CAG Gln	GTG Val	GTG Val	Asp	GTC Val 465	AAT Asn	GAC Asp	AAC Asn	GCA Ala	CCT Pro 470	GTC Val	TTC Phe	ACT Thr	CAG Gln	AGT Ser 475	GTC Val	1922

ACT Thr	GAC	G GTC	GCC Ala 480	Phe	e CCG	GAA Glu	AAC Asn	AAC Asn 485	Lys	CCT	GGT	GAP Glu	GTG Val 490	. Ile	GCT Ala	1	1970
GAG Glu	ATC Ile	Thr 495	Ala	AGT Ser	GAT Asp	GCT Ala	GAC Asp 500	Ser	GGC Gly	TCT Ser	AAT Asn	GCT Ala 505	Glu	CTG Leu	GTT Val	2	2018
TAC Tyr	Ser 510	Leu	GAG Glu	Pro	GAG Glu	CCG Pro 515	GCT Ala	GCT Ala	AAG Lys	GGC Gly	CTC Leu 520	Phe	ACC Thr	ATC	TCA Ser	2	066
CCC Pro 525	Glu	ACT Thr	GGA Gly	GAG Glu	ATC Ile 530	CAG Gln	GTG Val	AAG Lys	ACA Thr	TCT Ser 535	CTG Leu	GAT Asp	CGG Arg	GAA Glu	CAG Gln 540	2	114
CGG Arg	GAG Glu	AGC Ser	TAT Tyr	GAG Glu 545	TTG Leu	AAG Lys	GTG Val	GTG Val	GCA Ala 550	GCT Ala	GAC Asp	CGG Arg	GCC	AGT Ser 555	CCT Pro	2	162
AGC Ser	CTC Leu	CAG Gln	GGC Gly 560	ACA Thr	GCC Ala	ACT Thr	GTC Val	CTT Leu 565	GTC Val	AAT Asn	GTG Val	CTG Leu	GAC Asp 570	TGC Cys	AAT Asn	2.	210
Asp	Asn	575	Pro	Lys	Phe	ATG Met	Leu 580	Ser	Gly	Tyr	Asn	Phe 585	Ser	Val	Met	2:	258
GIU	Asn 590	Met	Pro	Ala	Leu	AGT Ser 595	Pro	Val	Gly	Met	Val 600	Thr	Val	Ile	Asp	2:	306
605	Asp	Lys	GIĀ	Glu	Asn 610	GCC Ala	Gln	Val	Gln	Leu 615	Ser	Val	Glu	Gln	Asp 620	2:	354
ASI	GIY	Asp	Pne	625	IIe	CAG Gln	Asn	Gly	Thr 630	Gly	Thr	Ile	Leu	Ser 635	Ser	24	402
Leu	ser	Pne	640	Arg	Glu	CAA Gln	Gln	Ser 645	Thr	Tyr	Thr	Phe	Gln 650	Leu	Lys	24	150
Ala	Val	655	GIĀ	GIÀ	Val	CCA Pro	Pro 660	Arg	Ser	Ala	Tyr	Val 665	Gly	Val	Thr	24	198
116	670	Val	Leu	Asp	Glu	AAT Asn 675	Asp	Asn	Ala	Pro	Tyr 680	Ile	Thr	Ala	Pro	25	46
685	VPII	Inr	ser	nis	690	CTG Leu	Leu	Thr	Pro	Gln 695	Thr	Arg	Leu	Gly	Glu 700	25	94
ACG Thr	GTC Val	AGC Ser	GIN	GTG Val 705	GCA Ala	GCC Ala	GAG Glu	GAC Asp	TTT Phe 710	GAC Asp	TCT Ser	GGT Gly	GTC Val	AAT Asn 715	GCC Ala	26	42

GA0 Glu	G CTO	ATO	720	: Ser	ATT	GCA Ala	GGI	GGC Gly 725	/ Asn	CCI Pro	TAT	GG;	7 Let 730	Phe	CAG Gln	2690
AT1 Ile	GGC Gly	TCF Ser 735	His	TCA Ser	Gly	GCC Ala	Ile 740	Thr	CTG Leu	GAG Glu	AAG Lys	GAG Glu 745	Ile	GAG	CGG	2738
CGC Arg	CAC His 750	His	GGG Gly	CTA Leu	CAC His	CGC Arg 755	CTG Leu	GTG Val	GTG Val	AAG Lys	GTC Val 760	Ser	GAC Asp	CGC Arg	GGC	2786
AAG Lys 765	Pro	CCA Pro	CGC Arg	TAT	GGC Gly 770	ACA Thr	GCC Ala	TTG Leu	GTC Val	CAT His 775	CTT	TAT Tyr	GTC Val	AAT Asn	GAG Glu 780	2834
ACT Thr	CTG	GCC Ala	AAC Asn	CGC Arg 785	ACG Thr	CTG Leu	CTG Leu	GAG Glu	ACC Thr 790	CTC Leu	CTG Leu	GCC	CAC	AGC Ser 795	CTG Leu	2882
GAC Asp	ACG Thr	CCG Pro	CTG Leu 800	GAT Asp	ATT	GAC Asp	ATT Ile	GCT Ala 805	GGG Gly	GAT Asp	CCA Pro	GAA Glu	TAT Tyr 810	Glu	CGC Arg	2930
TCC Ser	AAG Lys	CAG Gln 815	CGT	GGC Gly	AAC Asn	ATT Ile	CTC Leu 820	TTT Phe	GGT Gly	GTG Val	GTG Val	GCT Ala 825	GGT Gly	GTG Val	GTG Val	2978
GCC Ala	GTG Val 830	GCC Ala	TTG Leu	CTC Leu	ATC Ile	GCC Ala 835	CTG Leu	GCG Ala	GTT Val	CTT Leu	GTG Val 840	CGC Arg	TAC Tyr	TGC Cys	AGA Arg	3026
845	Arg	GIU	ATE	Lys	Ser 850	GGT Gly	Tyr	Gln	Ala	Gly 855	Lys	Lys	Glu	Thr	Lys 860	3074
nap	reu	TYP	Ala	865	rys	CCC Pro	Ser	Gly	Lys 870	Ala	Ser	Lys	Gly	Asn 875	Lys	3122
AGC Ser	AAA Lys	GGC Gly	AAG Lys 880	AAG Lys	AGC Ser	AAG Lys	TCC Ser	CCA Pro 885	AAG Lys	CCC Pro	GTG Val	AAG Lys	CCA Pro 890	GTG Val	GAG Glu	3170
GAC Asp	GAG Glu	GAT Asp 895	GAG Glu	GCC Ala	GGG Gly	CTG Leu	CAG Gln 900	AAG Lys	TCC Ser	CTC Leu	AAG Lys	TTC Phe 905	AAC Asn	CTG Leu	ATG Met	3218
AGC Ser	GAT Asp 910	GCC Ala	CCT Pro	GGG Gly	GAC Asp	AGT Ser 915	CCC Pro	CGC Arg	ATC Ile	CAC His	CTG Leu 920	CCC Pro	CTC Leu	AAC Asn	TAC Tyr	3266
CCA Pro 925	CCA Pro	GGC Gly	AGC Ser	PIO	GAC Asp 930	CTG Leu	GGC Gly	CGC Arg	His	TAT Tyr 935	CGC Arg	TCT Ser	AAC Asn	TCC Ser	CCA Pro 940	3314
CTG Leu	CCT Pro	TCC Ser	TTG	CAG Gln 945	CTG Leu	CAG (Gln)	CCC Pro	Gin	TCA Ser 950	CCC Pro	TCA Ser	GCC Ala	TCC Ser	AAG Lys 955	AAG Lys	3362

												TTC Phe				3410
GGG Gly	GAC Asp	ACC Thr 975	ACG Thr	TCC Ser	ACG Thr	GGC Gly	TCT Ser 980	GAG Glu	CAG Gln	TAC Tyr	TCC Ser	GAC Asp 985	TAC Tyr	AGC Ser	TAC Tyr	3458
CGC	ACC Thr 990	AAC Asn	CCC	CCC Pro	AAA Lys	TAC Tyr 995	CCC Pro	AGC Ser	AAG Lys	CAG Gln	TTA Leu 100	CCT Pro O	CAC His	CGC Arg	CGC Arg	3506
GTC Val 100	Thr	TTC Phe	TCG Ser	GCC Ala	ACC Thr 101	Ser	CAG Gln	GCC Ala	CAG Gln	GAG Glu 101	Leu	CAG Gln	GAC Asp	CCA Pro	TCC Ser 1020	3554
CAG Gln	CAC His	AGT Ser	TAC Tyr	TAT Tyr 102	Asp	AGT Ser	GGC Gly	CTG Leu	GAG Glu 1030	Glu	TCT Ser	GAG Glu	ACG Thr	CCG Pro 103	Ser	3602
AGC Ser	AAG Lys	TCA Ser	TCC Ser 1040	Ser	GGG Gly	CCT Pro	CGA Arg	CTC Leu 104	Gly	CCC Pro	CTG Leu	GCC Ala	CTG Leu 1050	Pro	GAG Glu	3650
GAT Asp	CAC His	TAT Tyr 105	Glu	CGC Arg	ACC Thr	ACC Thr	CCT Pro 1060	Asp	GGC Gly	AGC Ser	ATA Ile	GGA Gly 1065	Glu	ATG Met	GAG Glu	3698
CAC His	CCC Pro 1070	Glu	AAT Asn	GAC Asp	CTT Leu	CGC Arg 1075	Pro	TTG Leu	CCT Pro	GAT Asp	GTC Val 1080	GCC Ala	ATG Met	ACA Thr	GGC Gly	3746
ACA Thr 108	Сув	ACC Thr	CGG Arg	GAG Glu	TGC Cys 1090	Ser	GAG Glu	TTT Phe	GGC	CAC His 1095	Ser	GAC Asp	ACA Thr	TGC Cys	TGG Trp 1100	3794
ATG Met	CCT Pro	GGC Gly	CAG Gln	TCA Ser 1109	Ser	CCC Pro	AGC Ser	CGC Arg	CGG Arg 1110	Thr	AAG Lys	AGC Ser	AGC Ser	GCC Ala 1115	Leu	3842
AAA Lys	CTC Leu	TCC Ser	ACC Thr 1120	Phe	ATG Met	CCT Pro	TAC Tyr	CAG Gln 1125	Asp	CGA Arg	GGA Gly	GGG Gly	CAG Gln 1130	Glu	CCT Pro	3890
GCG Ala	GGC Gly	GCC Ala 1135	Gly	AGC Ser	CCC Pro	AGC Ser	CCC Pro 1140	Pro	GAA Glu	GAC Asp	CGG Arg	AAC Asn 1145	Thr	AAA Lys	ACG Thr	3938
GCC Ala	CCC Pro 1150	Val	CGC Arg	CTC Leu	CTG Leu	CCC Pro 1155	Ser	TAC Tyr	AGT Ser	GCC Ala	TTC Phe 1160	TCC Ser	CAC His	AGT Ser	AGC Ser	3986
CAT His 1165	Asp	TCC Ser	TGC Cys	AAG Lys	GAC Asp 1170	Ser	GCC Ala	ACC Thr	TTG Leu	GAG Glu 1175	Glu	ATC Ile	CCC Pro	CTG Leu	ACC Thr 1180	4034
CAG Gln	ACC Thr	TCG Ser	GAC Asp	TTC Phe 1185	Pro	CCC Pro	GCA Ala	GCC Ala	ACA Thr 1190	Pro	GCA Ala	TCT Ser	GCC Ala	CAG Gln 1195	Thr	4082

	GAG ATC TA G Glu Ile Ty 1200	AC CTG TGAGO yr Leu	CCCCT ACTG	CCCCC CCCCC	CTCCCC	4133
CAGCGCCGGC	CAGCTCCCAA	ATGCCCATTC	CAGGGCCTCA	CTCTCCACCC	CTTCAGCGTG	4193
GACTTCCTGC	CAGGGCCCAA	GTGGGGGTAT	CACTGACCTC	ATGACCACGC	TGGCCCTTCT	4253
CCCATGCAGG	GTCCAGGTCC	TCTCCCCTCA	TTTCCATCTC	CCAGCCCAGG	GGCCCCTTCC	4313
CCTTTATGGG	GCTTCCCCCA	GCTGATGCCC	AAGAGGGCTC	CTCTGCAATG	ACTGGGCTCC	4373
TTCCCTTGAC	TTCCAGGGAG	CACCCCTCG	ATTTGGGCAG	ATGGTGGAGT	CAAGGGTGGG	4433
CAGCGTACTT	CTAACTCATT	GTTTCCCTCA	TGGCCGACCA	GGGCGGGGAT	AGCATGCCCA	4493
ATTTTAGCCC	TGAAGCAGGG	CTGAACTGGG	GAGCCCCTTT	CCCTGGGAGC	TCCCAGAGGA	4553
AACTCTTGAC	CACCAGTGGC	TCCCTGAAGG	GCTTTTGTTA	CCAAAGGTGG	GGTAGGGACG	4613
GGGGTGGGAG	TGGAGCGGAG	GCCTTGTTTT	CCCGTGG			4650

(2) INFORMATION FOR SEQ ID NO:103:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1203 amino acids

 - (B) TYPE: amino acid(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:103:

Met Glu Pro Leu Arg His Ser Pro Gly Pro Gly Gly Gln Arg Leu Leu

Leu Pro Ser Met Leu Leu Ala Leu Leu Leu Leu Ala Pro Ser Pro

Gly His Ala Thr Arg Val Val Tyr Lys Val Pro Glu Glu Gln Pro Pro 35 40 45

Asn Thr Leu Ile Gly Ser Leu Ala Ala Asp Tyr Gly Phe Pro Asp Val 50 55 60

Gly His Leu Tyr Lys Leu Glu Val Gly Ala Pro Tyr Leu Arg Val Asp
65 75 80

Gly Lys Thr Gly Asp Ile Phe Thr Thr Glu Thr Ser Ile Asp Arg Glu 85 90 95

Gly Leu Arg Glu Cys Gln Asn Gln Leu Pro Gly Asp Pro Cys Ile Leu 100 105 110

Glu Phe Glu Val Ser Ile Thr Asp Leu Val Gln Asn Ala Ser Pro Arg

Leu Leu Glu Gly Gln Ile Glu Val Gln Asp Ile Asn Asp Asn Thr Pro Asn Phe Ala Ser Pro Val Ile Thr Leu Ala Ile Pro Glu Asn Thr Asn 150 Ile Gly Ser Leu Phe Pro Ile Pro Leu Ala Ser Asp Arg Asp Ala Gly 165 170 Pro Asn Gly Val Ala Ser Tyr Glu Leu Gln Val Ala Glu Asp Gln Glu Glu Lys Gln Pro Gln Leu Ile Val Met Gly Asn Leu Asp Arg Glu Arg 200 Trp Asp Ser Tyr Asp Leu Thr Ile Lys Val Gln Asp Gly Gly Ser Pro Pro Arg Ala Thr Ser Ala Leu Leu Arg Val Thr Val Leu Asp Thr Asn Asp Asn Ala Pro Lys Phe Glu Arg Pro Ser Tyr Glu Ala Glu Leu Ser Glu Asn Ser Pro Ile Gly His Ser Val Ile Gln Val Lys Ala Asn Asp Ser Asp Gln Gly Ala Asn Ala Glu Ile Glu Tyr Thr Phe His Gln Ala Pro Glu Val Val Arg Arg Leu Leu Arg Leu Asp Arg Asn Thr Gly Leu Ile Thr Val Gln Gly Pro Val Asp Arg Glu Asp Leu Ser Thr Leu Arg 315 Phe Ser Val Leu Ala Lys Asp Arg Gly Thr Asn Pro Lys Ser Ala Arg Ala Gln Val Val Val Thr Val Lys Asp Met Asn Asp Asn Ala Pro Thr Ile Glu Ile Arg Gly Ile Gly Leu Val Thr His Gln Asp Gly Met Ala Asn Ile Ser Glu Asp Val Ala Glu Glu Thr Ala Val Ala Leu Val Gln Val Ser Asp Arg Asp Glu Gly Glu Asn Ala Ala Val Thr Cys Val Val Ala Gly Asp Val Pro Phe Gln Leu Arg Gln Ala Ser Glu Thr Gly Ser 410 Asp Ser Lys Lys Lys Tyr Phe Leu Gln Thr Thr Thr Pro Leu Asp Tyr 425 Glu Lys Val Lys Asp Tyr Thr Ile Glu Ile Val Ala Val Asp Ser Gly

440

Asn Pro Pro Leu Ser Ser Thr Asn Ser Leu Lys Val Gln Val Val Asp Val Asn Asp Asn Ala Pro Val Phe Thr Gln Ser Val Thr Glu Val Ala 475 470 Phe Pro Glu Asn Asn Lys Pro Gly Glu Val Ile Ala Glu Ile Thr Ala Ser Asp Ala Asp Ser Gly Ser Asn Ala Glu Leu Val Tyr Ser Leu Glu Pro Glu Pro Ala Ala Lys Gly Leu Phe Thr Ile Ser Pro Glu Thr Gly Glu Ile Gln Val Lys Thr Ser Leu Asp Arg Glu Gln Arg Glu Ser Tyr 535 Glu Leu Lys Val Val Ala Ala Asp Arg Gly Ser Pro Ser Leu Gln Gly Thr Ala Thr Val Leu Val Asn Val Leu Asp Cys Asn Asp Asn Asp Pro Lys Phe Met Leu Ser Gly Tyr Asn Phe Ser Val Met Glu Asn Met Pro 585 Ala Leu Ser Pro Val Gly Met Val Thr Val Ile Asp Gly Asp Lys Gly Glu Asn Ala Gln Val Gln Leu Ser Val Glu Gln Asp Asn Gly Asp Phe Val Ile Gln Asn Gly Thr Gly Thr Ile Leu Ser Ser Leu Ser Phe Asp 630 635 Arg Glu Gln Gln Ser Thr Tyr Thr Phe Gln Leu Lys Ala Val Asp Gly Gly Val Pro Pro Arg Ser Ala Tyr Val Gly Val Thr Ile Asn Val Leu Asp Glu Asn Asp Asn Ala Pro Tyr Ile Thr Ala Pro Ser Asn Thr Ser 680 His Lys Leu Leu Thr Pro Gln Thr Arg Leu Gly Glu Thr Val Ser Gln Val Ala Ala Glu Asp Phe Asp Ser Gly Val Asn Ala Glu Leu Ile Tyr Ser Ile Ala Gly Gly Asn Pro Tyr Gly Leu Phe Gln Ile Gly Ser His Ser Gly Ala Ile Thr Leu Glu Lys Glu Ile Glu Arg Arg His His Gly Leu His Arg Leu Val Val Lys Val Ser Asp Arg Gly Lys Pro Pro Arg 755 760 765 Tyr Gly Thr Ala Leu Val His Leu Tyr Val Asn Glu Thr Leu Ala Asn
770 780

Arg Thr Leu Leu Glu Thr Leu Leu Gly His Ser Leu Asn Thr Pro Leu

Arg Thr Leu Leu Glu Thr Leu Leu Gly His Ser Leu Asp Thr Pro Leu 785 790 795 800

Asp Ile Asp Ile Ala Gly Asp Pro Glu Tyr Glu Arg Ser Lys Gln Arg 805 810 815

Gly Asn Ile Leu Phe Gly Val Val Ala Gly Val Val Ala Val Ala Leu 820 825 830

Leu Ile Ala Leu Ala Val Leu Val Arg Tyr Cys Arg Gln Arg Glu Ala 835 840 845

Lys Ser Gly Tyr Gln Ala Gly Lys Lys Glu Thr Lys Asp Leu Tyr Ala 850 855 860

Pro Lys Pro Ser Gly Lys Ala Ser Lys Gly Asn Lys Ser Lys Gly Lys 865 870 875 880

Lys Ser Lys Ser Pro Lys Pro Val Lys Pro Val Glu Asp Glu Asp Glu 885 890 895

Ala Gly Leu Gln Lys Ser Leu Lys Phe Asn Leu Met Ser Asp Ala Pro 900 905 910

Gly Asp Ser Pro Arg Ile His Leu Pro Leu Asn Tyr Pro Pro Gly Ser 915 920 925

Pro Asp Leu Gly Arg His Tyr Arg Ser Asn Ser Pro Leu Pro Ser Ile 930 935 940

Gln Leu Gln Pro Gln Ser Pro Ser Ala Ser Lys Lys His Gln Val Val 945 950 955 960

Gln Asp Leu Pro Pro Ala Asn Thr Phe Val Gly Thr Gly Asp Thr Thr 965 970 975

Ser Thr Gly Ser Glu Gln Tyr Ser Asp Tyr Ser Tyr Arg Thr Asn Pro 980 985 990

Pro Lys Tyr Pro Ser Lys Gln Leu Pro His Arg Arg Val Thr Phe Ser 995 1000 1005

Ala Thr Ser Gln Ala Gln Glu Leu Gln Asp Pro Ser Gln His Ser Tyr 1010 1015 1020

Tyr Asp Ser Gly Leu Glu Glu Ser Glu Thr Pro Ser Ser Lys Ser Ser 1025 1030 1035 1040

Ser Gly Pro Arg Leu Gly Pro Leu Ala Leu Pro Glu Asp His Tyr Glu 1045 1050 1055

Arg Thr Thr Pro Asp Gly Ser Ile Gly Glu Met Glu His Pro Glu Asn 1060 1065 1070

Asp Leu Arg Pro Leu Pro Asp Val Ala Met Thr Gly Thr Cys Thr Arg 1075 1080 1085

Glu	Сув 1090		Glu	Phe	Gly	His 1099	Ser	Asp	Thr	Cys	Trp 1100	Met)	Pro	Gly	Gln
Ser 1109		Pro	Ser	Arg	Arg 1110		Lys	Ser	Ser	Ala 111	Leu 5	Lys	Leu	Ser	Thr 1120
Phe	Met	Pro	Tyr	Gln 112		Arg	Gly	Gly	Gln 1130	Glu)	Pro	Ala	Gly	Ala 1135	Gly
Ser	Pro	Ser	Pro 1140		Glu	Asp	Arg	Asn 1145	Thr	Lys	Thr	Ala	Pro 1150	Val	Arg
Leu	Leu	Pro 1155		Tyr	Ser	Ala	Phe 1160	Ser	His	Ser	Ser	His 1169	Asp	Ser	Сув
Lys	Asp 1170		Ala	Thr	Leu	Glu 1179	Glu 5	Ile	Pro	Leu	Thr 1180	Gln)	Thr	Ser	Asp
Phe 118		Pro	Ala	Ala	Thr 1190		Ala	Ser	Ala	Gln 119		Ala	Lys	Arg	Glu 1200
Ile	Tyr	Leu													•
(2)	INFO	ORMA:	NOI	FOR	SEQ	ID 1	NO:10	04:							
	, – <i>,</i>	() () ()	A) LE B) TY C) ST O) TO	ENGTI PE: PRANI POLO	i: 27	789 leic ESS: line		pai:	cs						

- (ix) FEATURE:

 - (A) NAME/KEY: CDS (B) LOCATION: 115..2622
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:104:

CGAAAGCCAT G	TCGGACTCG TCGCCCA	GCG CCCAAGCGCT A	ACCCGCTGA AAGTT	TCTCA 60
GCGAAATCTC A	AGGGACGATC TGGACCC	CGC TGAGAGGAAC T	CCTTTTGAG TGAG	ATG 117 Met 1
	GCC TGG AGG AGC GC Ala Trp Arg Ser G: 5			
	CTT CTG CTT GGT GG Leu Leu Leu Gly A			_
	ATC CCG GAG GAA AG Ile Pro Glu Glu Ag 40			

	GTC Val															309
	CCG Pro															357
	ACC Thr															405
	GGG Gly															453
AAC Asn	CCG Pro 115	CTG Leu	GAG Glu	CTG Leu	TTC Phe	AGC Ser 120	GTG Val	GAA Glu	GTG Val	GTG Val	ATC Ile 125	CAG Gln	GAC Asp	ATC Ile	AAC Asn	501
	AAC Asn															549
GAG Glu	GCC Ala	GTG Val	GCT Ala	CCG Pro 150	GGG Gly	ACG Thr	CGC Arg	TTT Phe	CCG Pro 155	CTC Leu	GAG Glu	AGC Ser	GCG Ala	CAC His 160	GAT Asp	597
CCC Pro	GAT Asp	CTG Leu	GGA Gly 165	AGC Ser	AAC Asn	TCT Ser	TTA Leu	CAA Gln 170	ACC Thr	TAT Tyr	GAG Glu	CTG Leu	AGC Ser 175	CGA Arg	AAT Asn	645
GAA Glu	TAC Tyr	TTT Phe 180	GCG Ala	CTT Leu	CGC Arg	GTG Val	CAG Gln 185	ACG Thr	CGG Arg	GAG Glu	GAC Asp	AGC Ser 190	ACC Thr	AAG Lys	TAC Tyr	693
GCG Ala	GAG Glu 195	CTG Leu	GTG Val	TTG Leu	GAG Glu	CGC Arg 200	GCC Ala	CTG Leu	GAC Asp	CGA Arg	GAA Glu 205	CGG Arg	GAG Glu	CCT Pro	AGT Ser	741
CTC Leu 210	CAG Gln	TTA Leu	GTG Val	CTG Leu	ACG Thr 215	GCG Ala	TTG Leu	GAC Asp	GGA Gly	GGG Gly 220	ACC Thr	CCA Pro	GCT Ala	CTC Leu	TCC Ser 225	789
GCC Ala	AGC Ser	CTG Leu	CCT Pro	ATT Ile 230	CAC His	ATC Ile	AAG Lys	GTG Val	CTG Leu 235	GAC Asp	GCG Ala	AAT Asn	GAC Asp	AAT Asn 240	GCG Ala	837
CCT Pro	GTC Val	TTC Phe	AAC Asn 245	CAG Gln	TCC Ser	TTG Leu	TAC Tyr	CGG Arg 250	GCG Ala	CGC Arg	GTT Val	CCT Pro	GGA Gly 255	GGA Gly	TGC Cys	885
ACC Thr	TCC Ser	GGC Gly 260	ACG Thr	CGC Arg	GTG Val	GTA Val	CAA Gln 265	GTC Val	CTT Leu	GCA Ala	ACG Thr	GAT Asp 270	CTG Leu	GAT Asp	GAA Glu	933
GGC Gly	CCC Pro 275	AAC Asn	GGT Gly	GAA Glu	ATT Ile	ATT Ile 280	TAC Tyr	TCC Ser	TTC Phe	GGC Gly	AGC Ser 285	CAC His	AAC Asn	CGC Arg	GCC Ala	981

						GCC Ala										1029
						TTC Phe										1077
						GGC Gly										1125
						GAT Asp										1173
						CCA Pro 360										1221
						ACT Thr										1269
						CCG Pro										1317
CTC Leu	AAG Lys	AAT Asn	TAC Tyr 405	TTC Phe	ACT Thr	TTG Leu	AAA Lys	ACC Thr 410	AGT Ser	GCA Ala	GAC Asp	CTG Leu	GAT Asp 415	CGG Arg	GAG Glu	1365
ACT Thr	GTG Val	CCA Pro 420	GAA Glu	TAC Tyr	AAC Asn	CTC Leu	AGC Ser 425	ATC Ile	ACC Thr	GCC Ala	CGA Arg	GAC Asp 430	GCC Ala	GGA Gly	ACC Thr	1413
CCT Pro	TCC Ser 435	CTC Leu	TCA Ser	GCC Ala	CTT Leu	ACA Thr 440	ATA Ile	GTG Val	CGT Arg	GTT Val	CAA Gln 445	GTG Val	TCC Ser	GAC Asp	ATC Ile	1461
AAT Asn 450	GAC Asp	AAC Asn	CCT Pro	CCA Pro	CAA Gln 455	TCT Ser	TCT Ser	CAA Gln	TCT Ser	TCC Ser 460	TAC Tyr	GAC Asp	GTT Val	TAC Tyr	ATT Ile 465	1509
GAA Glu	GAA Glu	AAC Asn	AAC Asn	CTC Leu 470	CCC Pro	GGG Gly	GCT Ala	CCA Pro	ATA Ile 475	CTA Leu	AAC Asn	CTA Leu	AGT Ser	GTC Val 480	TGG Trp	1557
GAC Asp	CCC Pro	GAC Asp	GCC Ala 485	CCG Pro	CAG Gln	AAT Asn	GCT Ala	CGG Arg 490	CTT Leu	TCT Ser	TTC Phe	TTT Phe	CTC Leu 495	TTG Leu	GAG Glu	1605
CAA Gln	GGA Gly	GCT Ala 500	GAA Glu	ACC Thr	GGG Gly	CTA Leu	GTG Val 505	GGT Gly	CGC Arg	TAT Tyr	TTC Phe	ACA Thr 510	ATA Ile	AAT Asn	CGT Arg	1653
GAC Asp	AAT Asn 515	GGC Gly	ATA Ile	GTG Val	TCA Ser	TCC Ser 520	TTA Leu	GTG Val	CCC Pro	CTA Leu	GAC Asp 525	TAT Tyr	GAG Glu	GAT Asp	CGG Arg	1701

	Glu				ACA Thr 535											1749
					AGC Ser											1797
					CTA Leu											1845
					ACC Thr											1893
					GCA Ala											1941
					CAG Gln 615											1989
					CGT Arg											2037
					ATC Ile											2085
					GTG Val											2133
GCC Ala	GAG Glu 675	TTC Phe	CCC Pro	TCT Ser	GGC	TCT Ser 680	GCC Ala	ccc Pro	CGG Arg	GAG Glu	CAG Gln 685	AAA Lys	AAA Lys	AAT Asn	CTC Leu	2181
ACC Thr 690	TTT Phe	TAT Tyr	CTA Leu	CTT Leu	CTT Leu 695	TCT Ser	CTA Leu	ATC Ile	CTG Leu	GTT Val 700	TCT Ser	GTG Val	GGC Gly	TTC Phe	GTG Val 705	2229
GTC Val	ACA Thr	GTG Val	TTC Phe	GGA Gly 710	GTA Val	ATC Ile	ATA Ile	TTC Phe	AAA Lys 715	GTT Val	TAC Tyr	AAG Lys	TGG Trp	AAG Lys 720	CAG Gln	2277
TCT Ser	AGA Arg	GAC Asp	CTA Leu 725	TAC Tyr	CGA Arg	GCC Ala	CCG Pro	GTG Val 730	AGC Ser	TCA Ser	CTG Leu	TAC Tyr	CGA Arg 735	ACA Thr	CCA Pro	2325
GGG Gly	CCC Pro	TCC Ser 740	TTG Leu	CAC His	GCG Ala	GAC Asp	GCC Ala 745	GTG Val	CGG Arg	GGA Gly	GGC Gly	CTG Leu 750	ATG Met	TCG Ser	CCG Pro	2373
CAC	CTT Leu 755	TAC Tyr	CAT His	CAG Gln	GTG Val	TAT Tyr 760	CTC Leu	ACC Thr	ACG Thr	GAC Asp	TCC Ser 765	CGC Arg	CGC Arg	AGC Ser	GAC Asp	2421

								GCC Ala								2469
AAC Asn	ACG Thr	CTG Leu	CGG Arg	AGC Ser 790	TGT Cys	GAT Asp	CCG Pro	GTG Val	TTC Phe 795	TAT Tyr	AGG Arg	CAG Gln	GTG Val	TTG Leu 800	GGT Gly	2517
								GTA Val 810								2565
								TAC Tyr								2613
	TTT Phe 835		TAG	TGAT	GAA (GATG:	TTTT(CC TO	GTG1	atgc:	A TTO	CACA	CTTT			2662
CAAC	CTGG	CTC :	rtcc:	raga:	C A	AGT:	TAGTO	cc:	TTG:	rgag	ATG	TGG	CT (CCA	GAGTGT	2722
GGTT	TGT	GT (CCA	rttc	G G	GGA	AGAT	A CT	rgac:	CAT	CTG	rgga	CT 1	AATTO	CACATC	2782
CTC	AGCG															2789

(2) INFORMATION FOR SEQ ID NO:105:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 836 amino acids
 - (B) TYPE: amino acid
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: protein
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:105:

Met Val Pro Glu Ala Trp Arg Ser Gly Leu Val Ser Thr Gly Arg Val 1 5 15

Val Gly Val Leu Leu Leu Gly Ala Leu Asn Lys Ala Ser Thr Val 20 25 30

Ile His Tyr Glu Ile Pro Glu Glu Arg Glu Lys Gly Phe Ala Val Gly 35 40 45

Asn Val Val Ala Asn Leu Gly Leu Asp Leu Gly Ser Leu Ser Ala Arg
50 55 60

Arg Phe Pro Val Val Ser Gly Ala Ser Arg Arg Phe Phe Glu Val Asn 65 70 75 80

Arg Glu Thr Gly Glu Met Phe Val Asn Asp Arg Leu Asp Arg Glu Glu 85 90 95

Leu Cys Gly Thr Leu Pro Ser Cys Thr Val Thr Leu Glu Leu Val Val 100 105 110

Glu Asn Pro Leu Glu Leu Phe Ser Val Glu Val Val Ile Gln Asp Ile 115 120 125

Asn Asp Asn Asn Pro Ala Phe Pro Thr Gln Glu Met Lys Leu Glu Ile 135 Ser Glu Ala Val Ala Pro Gly Thr Arg Phe Pro Leu Glu Ser Ala His Asp Pro Asp Leu Gly Ser Asn Ser Leu Gln Thr Tyr Glu Leu Ser Arg Asn Glu Tyr Phe Ala Leu Arg Val Gln Thr Arg Glu Asp Ser Thr Lys 185 Tyr Ala Glu Leu Val Leu Glu Arg Ala Leu Asp Arg Glu Arg Glu Pro Ser Leu Gln Leu Val Leu Thr Ala Leu Asp Gly Gly Thr Pro Ala Leu Ser Ala Ser Leu Pro Ile His Ile Lys Val Leu Asp Ala Asn Asp Asn Ala Pro Val Phe Asn Gln Ser Leu Tyr Arg Ala Arg Val Pro Gly Gly Cys Thr Ser Gly Thr Arg Val Val Gln Val Leu Ala Thr Asp Leu Asp Glu Gly Pro Asn Gly Glu Ile Ile Tyr Ser Phe Gly Ser His Asn Arg Ala Gly Val Arg Gln Leu Phe Ala Leu Asp Leu Val Thr Gly Met Leu Thr Ile Lys Gly Arg Leu Asp Phe Glu Asp Thr Lys Leu His Glu Ile Tyr Ile Gin Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys Lys Val Leu Val Glu Val Val Asp Val Asn Asp Asn Ala Pro Glu Ile 345 Thr Val Thr Ser Val Tyr Ser Pro Val Pro Glu Asp Ala Ser Gly Thr Val Ile Ala Leu Leu Ser Val Thr Asp Leu Asp Ala Gly Glu Asn Gly Leu Val Thr Cys Glu Val Pro Pro Gly Leu Pro Phe Ser Leu Thr Ser 390 Ser Leu Lys Asn Tyr Phe Thr Leu Lys Thr Ser Ala Asp Leu Asp Arg Glu Thr Val Pro Glu Tyr Asn Leu Ser Ile Thr Ala Arg Asp Ala Gly Thr Pro Ser Leu Ser Ala Leu Thr Ile Val Arg Val Gln Val Ser Asp 440

Ile Asn Asp Asn Pro Pro Gln Ser Ser Gln Ser Ser Tyr Asp Val Tyr Ile Glu Glu Asn Asn Leu Pro Gly Ala Pro Ile Leu Asn Leu Ser Val Trp Asp Pro Asp Ala Pro Gln Asn Ala Arg Leu Ser Phe Phe Leu Leu Glu Gln Gly Ala Glu Thr Gly Leu Val Gly Arg Tyr Phe Thr Ile Asn Arg Asp Asn Gly Ile Val Ser Ser Leu Val Pro Leu Asp Tyr Glu Asp Arg Arg Glu Phe Glu Leu Thr Ala His Ile Ser Asp Gly Gly Thr Pro Val Leu Ala Thr Asn Ile Ser Val Asn Ile Phe Val Thr Asp Arg Asn Asp Asn Ala Pro Gln Val Leu Tyr Pro Arg Pro Gly Gly Ser Ser Val Glu Met Leu Pro Arg Gly Thr Ser Ala Gly His Leu Val Ser Arg Val Val Gly Trp Asp Ala Asp Ala Gly His Asn Ala Trp Leu Ser Tyr Ser Leu Phe Gly Ser Pro Asn Gln Ser Leu Phe Ala Ile Gly Leu His Thr 615 Gly Gln Ile Ser Thr Ala Arg Pro Val Gln Asp Thr Asp Ser Pro Arg Gln Thr Leu Thr Val Leu Ile Lys Asp Asn Gly Glu Pro Ser Leu Ser Thr Thr Ala Thr Leu Thr Val Ser Val Thr Glu Asp Ser Pro Glu Ala 665 Arg Ala Glu Phe Pro Ser Gly Ser Ala Pro Arg Glu Gln Lys Lys Asn Leu Thr Phe Tyr Leu Leu Leu Ser Leu Ile Leu Val Ser Val Gly Phe Val Val Thr Val Phe Gly Val Ile Ile Phe Lys Val Tyr Lys Trp Lys 720 Gln Ser Arg Asp Leu Tyr Arg Ala Pro Val Ser Ser Leu Tyr Arg Thr Pro Gly Pro Ser Leu His Ala Asp Ala Val Arg Gly Gly Leu Met Ser Pro His Leu Tyr His Gln Val Tyr Leu Thr Thr Asp Ser Arg Arg Ser

								-	107	•					
Asp	Pro 770	Leu	Leu	Lys	Lys	Pro 775	Gly	Ala	Ala	Ser	Pro 780	Leu	Ala	Ser	Arg
Gln 785	Asn	Thr	Leu	Arg	Ser 790	Сув	Asp	Pro	Val	Ph e 795	Tyr	Arg	Gln	Val	Leu 800
Gly	Ala	Glu	Ser	Ala 805	Pro	Pro	Gly	Gln	Val 810	Arg	Phe	Ser	Lys	Ser 815	Сув
Leu	Thr	Leu	Leu 820	Val	Pro	Phe	Tyr	Ser 825	Tyr	Ile	Ile	Leu	Arg 830	Arg	Leu
Glu	Leu	Phe 835	Phe												
(2)	INFO	RMAI	CION	FOR	SEQ	ID N	10:10	06:							
	(i)	(F (E	QUENC A) LE B) TY C) SI O) TO	NGTI PE: RANI	nucl	751 E Leic ESS:	ase ació sino	pair 1	: s						
	(ii)	MOI	ECUI	E TY	PE:	CDNA	1								

- (ix) FEATURE:
 (A) NAME/KEY: CDS
 (B) LOCATION: 115..2160
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:106:

CGAAAGCCAT GTCGGACTCG TCGCCCAGCG CCCAAGCGCT AACCCGCTGA AAGTTTCTCA 6										A 60						
GCG.	AAAT(CTC :	AGGG	ACGA:	IC T	GGAC	CCCG	C TG	AGAG	GAAC	TGC	TTTT(GAG 1	TGAG	ATG Met 1	117
GTC Val	CCA Pro	GAG Glu	GCC Ala 5	TGG Trp	AGG Arg	AGC Ser	GGA Gly	CTG Leu 10	GTA Val	AGC Ser	ACC Thr	GLY	AGG Arg 15	GTA Val	GTG Val	165
GGA Gly	GTT Val	TTG Leu 20	CTT Leu	CTG Leu	CTT Leu	GGT Gly	GCC Ala 25	TTG Leu	AAC Asn	AAG Lys	GCT Ala	TCC Ser 30	ACG Thr	GTC Val	ATT Ile	213
CAC His	TAT Tyr 35	GAG Glu	ATC Ile	CCG Pro	GAG Glu	GAA Glu 40	AGA Arg	GAG Glu	AAG Lys	GGT Gly	TTC Phe 45	GCT Ala	GTG Val	G17 GCC	AAC Asn	261
GTG Val 50	GTC Val	GCG Ala	AAC Asn	CTT Leu	GGT Gly 55	TTG Leu	GAT Asp	CTC Leu	GGT Gly	AGC Ser 60	CTC Leu	TCA Ser	GCC Ala	CGC Arg	AGG Arg 65	309
TTC Phe	CCG Pro	GTG Val	GTG Val	TCT Ser 70	GGA Gly	GCT Ala	AGC Ser	CGA Arg	AGA Arg 75	TTC Phe	TTT Phe	GAG Glu	GTG Val	AAC Asn 80	CGG Arg	357

	ACC Thr															405
	GGG Gly															453
	CCG Pro 115															501
	AAC Asn															549
	GCC Ala															597
	GAT Asp															645
Glu	TAC Tyr	Phe 180	Ala	Leu	Arg	Val	Gln 185	Thr	Arg	Glu	Asp	Ser 190	Thr	Lys	Tyr	693
	GAG Glu 195															741
Leu 210	CAG Gln	Leu	Val	Leu	Thr 215	Ala	Leu	Asp	Gly	Gly 220	Thr	Pro	Ala	Leu	Ser 225	789
GCC Ala	AGC Ser	CTG Leu	CCT Pro	ATT Ile 230	CAC His	ATC Ile	AAG Lys	GTG Val	CTG Leu 235	GAC Asp	GCG Ala	AAT Asn	GAC Asp	AAT Asn 240	GCG Ala	837
CCT Pro	GTC Val	TTC Phe	AAC Asn 245	CAG Gln	TCC Ser	TTG Leu	TAC Tyr	CGG Arg 250	GCG Ala	CGC Arg	GTT Val	CCT Pro	GGA Gly 255	GGA Gly	TGC Cys	885
ACC Thr	TCC Ser	GGC Gly 260	ACG Thr	CGC Arg	GTG Val	GTA Val	CAA Gln 265	GTC Val	CTT Leu	GCA Ala	ACG Thr	GAT Asp 270	CTG Leu	GAT Asp	GAA Glu	933
GGC Gly	Pro 275	AAC Asn	GGT Gly	GAA Glu	ATT Ile	ATT Ile 280	TAC Tyr	TCC Ser	TTC Phe	GGC Gly	AGC Ser 285	CAC His	AAC Asn	CGC Arg	GCC Ala	981
GGC Gly 290	GTG Val	CGG Arg	CAA Gln	CTA Leu	TTC Phe 295	GCC Ala	TTA Leu	GAC Asp	CTT Leu	GTA Val 300	ACC Thr	GGG Gly	ATG Met	CTG Leu	ACA Thr 305	1029
ATC Ile	AAG Lys	GGT Gly	CGG Arg	CTG Leu 310	GAC Asp	TTC Phe	GAG Glu	GAC Asp	ACC Thr 315	AAA Lys	CTC Leu	CAT His	GAG Glu	ATT Ile 320	TAC Tyr	1077

ATC Ile	CAG Gln	GCC Ala	AAA Lys 325	GAC Asp	AAG Lys	GGC Gly	GCC Ala	AAT Asn 330	CCC Pro	GAA Glu	GGA Gly	GCA Ala	CAT His 335	TGC Cys	AAA Lys		1125
GTG Val	TTG Leu	GTG Val 340	Glu	GTT Val	GTG Val	GAT Asp	GTG Val 345	AAT Asn	GAC Asp	AAC Asn	GCC Ala	CCG Pro 350	GAG Glu	ATC Ile	ACA Thr		1173
GTC Val	ACC Thr 355	TCC Ser	GTG Val	TAC Tyr	AGC Ser	CCA Pro 360	GTA Val	CCC Pro	GAG Glu	GAT Asp	GCC Ala 365	TCT Ser	GGG Gly	ACT Thr	GTC Val		1221
ATC Ile 370	Ala	TTG Leu	CTC Leu	AGT Ser	GTG Val 375	ACT Thr	GAC Asp	CTG Leu	GAT Asp	GCT Ala 380	GGC Gly	GAG Glu	AAC Asn	GGG Gly	CTG Leu 385		1269
GTG Val	ACC Thr	TGC Cys	GAA Glu	GTT Val 390	CCA Pro	CCG Pro	GGT Gly	CTC Leu	CCT Pro 395	TTC Phe	AGC Ser	CTT Leu	ACT Thr	TCT Ser 400	TCC Ser		1317
CTC Leu	AAG Lys	AAT Asn	TAC Tyr 405	TTC Phe	ACT Thr	TTG Leu	AAA Lys	ACC Thr 410	AGT Ser	GCA Ala	GAC Asp	CTG Leu	GAT Asp 415	CGG Arg	GAG Glu		1365
ACT Thr	GTG Val	CCA Pro 420	GAA Glu	TAC Tyr	AAC Asn	CTC Leu	AGC Ser 425	ATC Ile	ACC Thr	GCC Ala	CGA Arg	GAC Asp 430	GCC Ala	GGA Gly	ACC Thr	,	1413
CCT Pro	TCC Ser 435	CTC Leu	TCA Ser	GCC Ala	CTT Leu	ACA Thr 440	ATA Ile	GTG Val	CGT Arg	GTT Val	CAA Gln 445	GTG Val	TCC Ser	GAC Asp	ATC Ile	;	1461
AAT Asn 450	GAC Asp	AAC Asn	CCT Pro	CCA Pro	CAA Gln 455	TCT Ser	TCT Ser	CAA Gln	TCT Ser	TCC Ser 460	TAC Tyr	GAC Asp	GTT Val	TAC Tyr	ATT Ile 465	:	1509
GAA Glu	GAA Glu	AAC Asn	AAC Asn	CTC Leu 470	CCC Pro	GGG Gly	GCT Ala	CCA Pro	ATA Ile 475	CTA Leu	AAC Asn	CTA Leu	AGT Ser	GTC Val 480	TGG Trp	:	1557
GAC Asp	CCC Pro	GAC Asp	GCC Ala 485	CCG Pro	CAG Gln	AAT Asn	GCT Ala	CGG Arg 490	CTT Leu	TCT Ser	TTC Phe	TTT Phe	CTC Leu 495	TTG Leu	GAG Glu	:	1605
CAA Gln	GGA Gly	GCT Ala 500	GAA Glu	ACC Thr	GGG Gly	CTA Leu	GTG Val 505	GGT Gly	CGC Arg	TAT Tyr	TTC Phe	ACA Thr 510	ATA Ile	AAT Asn	CGT Arg	:	1653
GAC Asp	AAT Asn 515	GGC Gly	ATA Ile	GTG Val	TCA Ser	TCC Ser 520	TTA Leu	GTG Val	CCC Pro	CTA Leu	GAC Asp 525	TAT Tyr	GAG Glu	GAT Asp	CGG Arg	1	1701
CGG Arg 530	GAA Glu	TTT Phe	GAA Glu	TTA Leu	ACA Thr 535	GCT Ala	CAT His	ATC Ile	AGC Ser	GAT Asp 540	GGG	GLY	ACC Thr	CCG Pro	GTC Val 545	1	1749
CTA Leu	GCC Ala	ACC Thr	AAC Asn	ATC Ile 550	AGC Ser	GTG Val	AAC Asn	ATA Ile	TTT Phe 555	GTC Val	ACT Thr	GAT Asp	CGC Arg	AAT Asn 560	GAC Asp	1	1797

AAT Asn	GCC Ala	CCC Pro	CAG Gln 565	GTC Val	CTA Leu	TAT Tyr	CCT Pro	CGG Arg 570	CCA Pro	GGT Gly	GLY	AGC Ser	TCG Ser 575	GTG Val	GAG Glu		1845
ATG Met	CTG Leu	CCT Pro 580	CGA Arg	GGT Gly	ACC Thr	TCA Ser	GCT Ala 585	GGC Gly	CAC His	CTA Leu	GTG Val	TCA Ser 590	CGG Arg	GTG Val	GTA Val		1893
GGC Gly	TGG Trp 595	GAC Asp	GCG Ala	GAT Asp	GCA Ala	GGG Gly 600	CAC His	AAT Asn	GCC Ala	TGG Trp	CTC Leu 605	TCC Ser	TAC Tyr	AGT Ser	CTC Leu		1941
TTT Phe 610	GGA Gly	TCC Ser	CCT Pro	AAC Asn	CAG Gln 615	AGC Ser	CTT Leu	TTT Phe	GCC Ala	ATA Ile 620	GGG	CTG Leu	CAC His	ACT Thr	GGT Gly 625		1989
CAA Gln	ATC Ile	AGT Ser	ACT Thr	GCC Ala 630	CGT Arg	CCA Pro	GTC Val	CAA Gln	GAC Asp 635	ACA Thr	GAT Asp	TCA Ser	CCC Pro	AGG Arg 640	CAG Gln		2037
ACT Thr	CTC Leu	ACT Thr	GTC Val 645	TTG Leu	ATC Ile	AAA Lys	GAC Asp	AAT Asn 650	GGG Gly	GAG Glu	CCT Pro	TCG Ser	CTC Leu 655	TCC Ser	ACC Thr		2085
ACT Thr	GCT Ala	ACC Thr 660	CTC Leu	ACT Thr	GTG Val	TCA Ser	GTA Val 665	ACC Thr	GAG Glu	GAC Asp	TCT Ser	CCT Pro 670	GAA Glu	GCC Ala	CGA Arg		2133
GCC Ala	GAG Glu 675	TTC Phe	CCC Pro	TCT Ser	GGC Gly	TCT Ser 680	GCC Ala	AGT Ser	TAAA	CCTI	CT I	TAAT	TATG	G			2180
ATTA	GCCA	TT A	ACAT	TTTT	G AA	ACGT	GGAC	CAT	TTAA	CCT	CGGC	CTAC	cc c	CTCC	AACTG		2240
TCCI	GGTG	AT G	AGTT	CATT	'A GC	TAAG	TTAA	ATT	AATT	GAA	CTTI	GATO	TA A	ACCA	AAACA	:	2300
AATC	AGGA	AA A	AAAT	.GCTG	T AA	AGGA	ACTT	ATC	AAGC	ATT	CCAA	AACC	AA C	TAGA	AATTA	:	2360
CTTG	AAGT	TT C	GAGT	GAGC	A TT	GCCT	GTGC	CAG	TATT	CTT	CATT	ATAG	GA I	TATA	AACTC	;	2420
GTTT	TTTT	CC C	AAAG	CGCA	T GT	CTAC	GCCA	GGC	AGAG	GAG	TAAT	TATT	'CA G	CCAA	TTTCA	:	2480
TGGA	TGTA	AC G	ATGG	ATAT	A AA	TAAT	TGAT	AGC	ACCT	AGA	GGCT	TCCA	GT T	TGGG	TGGAA	:	2540
															TAGTT		2600
															ATTAG		2660
									AAAT	ATA	AAAA	GCCA	AA C	TTTA	AATAA	:	2720
ATCA	TAGA	GA C	CTCA	GACA	T AA	TATA	GGAA	A									2751

(2) INFORMATION FOR SEQ ID NO:107:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 682 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:107:

Met Val Pro Glu Ala Trp Arg Ser Gly Leu Val Ser Thr Gly Arg Val Val Gly Val Leu Leu Leu Gly Ala Leu Asn Lys Ala Ser Thr Val Ile His Tyr Glu Ile Pro Glu Glu Arg Glu Lys Gly Phe Ala Val Gly Asn Val Val Ala Asn Leu Gly Leu Asp Leu Gly Ser Leu Ser Ala Arg Arg Phe Pro Val Val Ser Gly Ala Ser Arg Arg Phe Phe Glu Val Asn Arg Glu Thr Gly Glu Met Phe Val Asn Asp Arg Leu Asp Arg Glu Glu 85 90 95 Leu Cys Gly Thr Leu Pro Ser Cys Thr Val Thr Leu Glu Leu Val Val Glu Asn Pro Leu Glu Leu Phe Ser Val Glu Val Val Ile Gln Asp Ile Asn Asp Asn Asn Pro Ala Phe Pro Thr Gln Glu Met Lys Leu Glu Ile Ser Glu Ala Val Ala Pro Gly Thr Arg Phe Pro Leu Glu Ser Ala His Asp Pro Asp Leu Gly Ser Asn Ser Leu Gln Thr Tyr Glu Leu Ser Arg Asn Glu Tyr Phe Ala Leu Arg Val Gln Thr Arg Glu Asp Ser Thr Lys Tyr Ala Glu Leu Val Leu Glu Arg Ala Leu Asp Arg Glu Arg Glu Pro Ser Leu Gln Leu Val Leu Thr Ala Leu Asp Gly Gly Thr Pro Ala Leu 210 215 220 Ser Ala Ser Leu Pro Ile His Ile Lys Val Leu Asp Ala Asn Asp Asn Ala Pro Val Phe Asn Gln Ser Leu Tyr Arg Ala Arg Val Pro Gly Gly Cys Thr Ser Gly Thr Arg Val Val Gln Val Leu Ala Thr Asp Leu Asp Glu Gly Pro Asn Gly Glu Ile Ile Tyr Ser Phe Gly Ser His Asn Arg Ala Gly Val Arg Gln Leu Phe Ala Leu Asp Leu Val Thr Gly Met Leu 295 Thr Ile Lys Gly Arg Leu Asp Phe Glu Asp Thr Lys Leu His Glu Ile

Tyr Ile Gln Ala Lys Asp Lys Gly Ala Asn Pro Glu Gly Ala His Cys Lys Val Leu Val Glu Val Val Asp Val Asn Asp Asn Ala Pro Glu Ile Thr Val Thr Ser Val Tyr Ser Pro Val Pro Glu Asp Ala Ser Gly Thr Val Ile Ala Leu Leu Ser Val Thr Asp Leu Asp Ala Gly Glu Asn Gly Leu Val Thr Cys Glu Val Pro Pro Gly Leu Pro Phe Ser Leu Thr Ser Ser Leu Lys Asn Tyr Phe Thr Leu Lys Thr Ser Ala Asp Leu Asp Arg 410 Glu Thr Val Pro Glu Tyr Asn Leu Ser Ile Thr Ala Arg Asp Ala Gly Thr Pro Ser Leu Ser Ala Leu Thr Ile Val Arg Val Gln Val Ser Asp Ile Asn Asp Asn Pro Pro Gln Ser Ser Gln Ser Ser Tyr Asp Val Tyr Ile Glu Glu Asn Asn Leu Pro Gly Ala Pro Ile Leu Asn Leu Ser Val Trp Asp Pro Asp Ala Pro Gln Asn Ala Arg Leu Ser Phe Phe Leu Leu Glu Gln Gly Ala Glu Thr Gly Leu Val Gly Arg Tyr Phe Thr Ile Asn 505 Arg Asp Asn Gly Ile Val Ser Ser Leu Val Pro Leu Asp Tyr Glu Asp 520 Arg Arg Glu Phe Glu Leu Thr Ala His Ile Ser Asp Gly Gly Thr Pro 535 Val Leu Ala Thr Asn Ile Ser Val Asn Ile Phe Val Thr Asp Arg Asn Asp Asn Ala Pro Gln Val Leu Tyr Pro Arg Pro Gly Gly Ser Ser Val Glu Met Leu Pro Arg Gly Thr Ser Ala Gly His Leu Val Ser Arg Val Val Gly Trp Asp Ala Asp Ala Gly His Asn Ala Trp Leu Ser Tyr Ser 600 Leu Phe Gly Ser Pro Asn Gln Ser Leu Phe Ala Ile Gly Leu His Thr 615 Gly Gln Ile Ser Thr Ala Arg Pro Val Gln Asp Thr Asp Ser Pro Arg

Gln Thr Leu Thr Val Leu Ile Lys Asp Asn Gly Glu Pro Ser Leu Ser 650

Thr Thr Ala Thr Leu Thr Val Ser Val Thr Glu Asp Ser Pro Glu Ala

Arg Ala Glu Phe Pro Ser Gly Ser Ala Ser

(2) INFORMATION FOR SEQ ID NO:108:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 2831 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: single

 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:108:

GAATTCGGCA	CGAGGCTGAA	CTGAGGGTGA	CGGACATAAA	CGACTATTCT	CCAGTGTTCA	60
GTGAAAGAGA	AATGATACTG	AGGATACCAG	AAAACAGTGC	TCGGGGAAAT	ACATTCCCTT	120
TAAACAATGC	TCTGGACTCA	GACGTAGATA	TCAACAATAT	CCAGACCTAT	AGGCTCAGCT	180
CAAACTCTCA	TTTCCTGGTT	GTAACCCGCA	ACCGCAGTGA	TGGCAGGAAG	TACCCAGAGC	240
TGGTGCTGGA	GAAAGAACTG	GATCGAGAGG	AGGAACCTGA	GCTGAGGTTA	ACGCTGACAG	300
CTTTGGATGG	TGGCTCTCCT	CCCCGGTCTG	GGACGACACA	GGTCCTCATT	GAAGTAGTGG	360
ACACCAACGA	TAATGCACCC	GAGTTTCAGC	AGCCAACATA	CCAAGTGCAA	ACTCCCGAGA	420
ACAGTCCCAC	CGGCTCTCTG	GTACTCACAG	TCTCAGCCAA	TGACTTAGAC	AGTGGAGACT	480
ATGGGAAAGT	CTTGTACGCA	CTTTCGCAAC	CCTCAGAAGA	TATTAGCAAA	ACATTCGAGG	540
TAAACCCTGT	AACCGGGGAA	ATTCGCCTAC	GAAAAGAGGT	GAATTTTGAA	ACTATTCCTT	600
CGTATGAAGT	GGTTATCAAG	GGGACGGACG	GGGGAGGTCT	CTCAGGAAAA	TGCACTCTGT	660
TACTGCAGGT	GGTGGACGTG	AATGACAATG	CCCCAGAAGT	GATGCTATCT	GCGCTAACCA	720
ACCCAGTCCC	AGAAAATTCC	CCCGATGAGG	TAGTGGCTGT	TTTCAGTGTT	AGAGATCCTG	780
ACTCTGGGAA	CAACGGAAAA	GTGATTGCAT	CCATCGAGGA	AGACCTGCCC	TTTCTTCTAA	840
AATCTTCAGG	AAAGAACTTT	TACACTTTAG	TAACCAAGGG	AGCACTTGAC	AGGGAAGAAA	900
GAGAGCAATT	GAACATCACC	ATCACAGTCA	CTGACCTGGG	CATACCCAGG	CTCACCACCC	960
AACACACCAT	AACAGTGCAG	GTGGCAGACA	TCAACGACAA	TGCCCCCTCC	TTCACCCAAA	1020
CCTCCTACAC	CATGTTTGTC	CGCGAGAACA	ACAGCCCCGC	CCTGCACATA	GGCACCATCA	1080
GCGCCACAGA	CTCAGACTCA	GGATCCAATG	CCCACATCAC	CTACTCGCTG	CTACCGCCCC	1140

AAGACCCACA	GCTGGCCCTC	GACTCGCTCA	TCTCCATCAA	TGTAGACAAC	GGGCAGCTGT	1200
TCGCGCTCAG	GGCGCTAGAC	TATGAGGCTC	TGCAGGGCTT	CGAGTTCCAT	GTGGGCGCCA	1260
CAGACCAAGG	CTCGCCCGCG	CTCAGCAGCC	AGGCTCTGGT	GCACGTGGTG	GTGTTGGACG	1320
ACAATGACAA	TGCGCCCTTC	GTGCTCTACC	CGCTGCAAAA	CGCCTCTGCA	CCCTTCACTG	1380
AGCTGCTGCC	CAGGGCGGCA	GAGCCTGGAT	ACCTGGTTAC	CAAGGTGGTA	GCTGTGGACC	1440
GCGACTCTGG	CCAGAATGCC	TGGCTGTCAT	TCCAGCTGCT	CAAGGCCACG	GAGCCCGGGC	1500
TGTTCAACGT	ATGGGCGCAC	AATGGCGAGG	TACGCACCTC	CAGGCTGCTG	AGCGAGCGCG	1560
ACGCACCCAA	GCACAAGCTG	CTGCTGTTGG	TCAAGGACAA	TGGAGATCCT	CCACGCTCTG	1620
CCAGTGTTAC	TCTGCACGTG	CTAGTGGTGG	ATGCCTTCTC	TCAGCCCTAC	CTGCCTCTGC	1680
CAGAGGTGGC	GCACGACCCT	GCACAAGAAG	AAGATGCGCT	AACACTCTAC	CTGGTCATAG	1740
CTTTGGCATC	TGTGTCTTCT	CTCTTCCTCT	TGTCTGTGCT	GCTGTTCGTG	GGGGTGAGGC	1800
TCTGCAGGAG	GGCCAGGGCA	GCCTCTCTGA	GTGCCTATTC	TGTGCCTGAA	GGCCACTTTC	1860
CTGGCCAGCT	GGTGGATGTC	AGAGGTATGG	GGACCCTGTC	CCAGAGCTAC	CAGTATGATG	1920
TATGTCTGAT	GGGGGATTCT	TCTGGGACCA	GCGAATTTAA	CTTCTTAAAG	CCAGTTCTGC	1980
CTAGCTCTCT	GCACCAGTGC	TCTGGGAAAG	AAATAGAGGA	AAATTCCACA	CTCCAGAATA	2040
GTTTTGGGTT	TCATCATTAA	TAGAAAACTA	CTTTACAGAT	ATTTAATTCC	AAATATCATC	2100
TTGTTGATTA	ACTAAAGTCT	GTTCACATGT	AGCTAGCTAG	CAACGATTTT	AATGTTCACT	2160
TTACCCATCT	TTTTTCAGGG	TCATGTCTAA	AGCTACAAGT	TTGNCTTTAC	TTATACTTGT	2220
CGCACAGAAT	NNNNNNNN	TGGTGTATAA	GTCACAGTCA	TGGGATACTG	GCACAAGATG	2280
GCAGCTTGAT	TGCTCAGTTA	TGGCTGCAAA	GGGGNGCTTG	AGTTTAGGGA	ATGTGTTAGA	2340
GCTGGAATAA	GTTTTCTGAG	AAATGTGTAA	GACAAATTTC	TTTTGCACAT	TCCCTGTGTT	2400
CCTGTACCCC	TGTTTCCAGA	ACTACGAAAT	GTGTCATCAG	AAGGCATGCT	CACATTTTCC	2460
CCTTTGTTTG	CGTGACCCGG	GTGCCAGAAA	TTAAATAAAA	TTAGCATGGA	GTTCAATGCA	2520
GCATTAAAAC	AAAGTTACTT	CTACAAACCT	TTTATTCGAC	GGTTAAAATT	GTAACTTCCC	2580
CACCCATGAG	GCTGGCTGTA	AGAACCAGTA	TGAATGGGTG	TCTATCGCAA	CCTTATTTTC	2640
АААААТСААА	CAAAAGGAGA	AATGAGAGAC	CAAACAACAC	GCTACAGGAA	AGATTTCATA	2700
AGGATGTATG	TATGGACACA	AAAACTGGGA	TACAGACATT	TTAAATCTGT	TGGTACCACA	2760
TGGTGGCGCT	GCAGGCTAAA	GAAATGCAAG	GGAAATTAAA	AAGAGGCTGA	GCTAGAAGTC	2820
АААААААА	A					2831

(2) INFORMATION FOR SEQ ID NO:109:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 3353 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA
- (ix) FEATURE:
 (A) NAME/KEY: CDS
 (B) LOCATION: 763..3123
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:109:

GTATTTTTCC ACAC	GTTTAAA ATTTTCATÄ	A AATCATAACT	CTCTGACTTT ATGTAGAAAG	60
GATACCACAC TGG	aattaac Gtgtagcti	T TTCTTGATGT	AATCCAACCA ATGGGAGCAC	120
AATTCTGGTA CATA	AGGCTGT CTAGAATTT	G AAAGAAATTA	AAGAATTCAT TTTGTTTTGC	180
TGATAAATTT TTAA	AGAAATC ACGTGGCTT	T ATGTTATTAT	TATTACAAGA TGACTGATCA	240
CTATTATGTC TTC	ITTCACT TCTCAATTT	C CCTCAGAACA	CTACACCCAG ACTACAGGCT	300
CTGGAGGGTG GGG	ACCATGT CTGGGTTGT	T TACTGATGTA	TTTCATAATT TGGCACATAG	360
AGACCAATAA TACT	iccitta aatgaagaa	A TTAATAATTA	CCATTGCGTG ATATTGTGAT	420
TACATCATTT CCTC	CCCAATT TCCAAACTC	C TAATAGAATA	GAGAATAGAT CAATTGTAGC	480
AATTCGTTTC GAAC	GCAAAGA CAACGCATG	G TGGCGCTGCA	GGCTAAGGCT TCAAAAAAAG	540
GAAAAGGAAA AAGC	CCCATGA AATGCTACT	A GCTACTTCAG	ACCTCTTTCA GCCTAAGAGG	600
AAAGCCTGTT AGCA	AGAGCAC GGACCAGTG	T CTCCGGAGAA	TGCTATTCTC CTACATTTCC	660
GAACAGGTTA TCAA	ACGCACA GATCGATCA	C TGCCTCTGTC	CCATCGCTCC CTGAAGTAGC	720
TCTGACTCCG GTTC	CCTTGAA AGGGGCGTG	T ACAGAAGTAA	AG ATG GAG CCT GCA Met Glu Pro Ala 1	774
GGG GAG CGC TTT Gly Glu Arg Phe 5	CCC GAA CAA AGG Pro Glu Gln Arg 10	CAA GTC CTG Gln Val Leu 15	ATT CTC CTT CTT TTA Ile Leu Leu Leu Leu 20	822
CTG GAA GTG ACT Leu Glu Val Thr	CTG GCA GGC TGG Leu Ala Gly Trp 25	GAA CCC CGT Glu Pro Arg 30	CGC TAT TCT GTG ATG Arg Tyr Ser Val Met 35	870
GAG GAA ACA GAG Glu Glu Thr Glu 40	ı Arg Gly Ser Phe	GTA GCC AAC Val Ala Asn 45	CTG GCC AAT GAC CTA Leu Ala Asn Asp Leu 50	918
GGG CTG GGA GTG Gly Leu Gly Val 55	GGG GAG CTA GCC Gly Glu Leu Ala 60	Glu Arg Gly	GCC CGG GTA GTT TCT Ala Arg Val Val Ser 65	966

		AAC Asn															1014
		AAT Asn															1062
		ATA Ile															1110
		GCT Ala															1158
		GAA Glu 135															1206
		GTG Val															1254
		GTT Val															1302
		CGC Arg															1350
		GAA Glu															1398
		GTG Val 215															1446
CTC Leu	ATC Ile 230	TTG Leu	GTC Val	TTG Leu	GAC Asp	GCC Ala 235	AAT Asn	GAC Asp	AAT Asn	GCC Ala	CCG Pro 240	GAG Glu	TTT Phe	GTG Val	CAG Gln		1494
GCG Ala 245	CTC Leu	TAC Tyr	GAG Glu	GTG Val	CAG Gln 250	GTC Val	CCA Pro	GAG Glu	AAC Asn	AGC Ser 255	CCA Pro	GTA Val	GGC Gly	TCC Ser	CTA Leu 260	•	1542
GTT Val	GTC Val	AAG Lys	GTC Val	TCT Ser 265	GCT Ala	AGG Arg	GAT Asp	TTA Leu	GAC Asp 270	ACT Thr	GGG Gly	ACA Thr	AAT Asn	GGA Gly 275	GAG Glu		1590
ATA Ile	TCA Ser	TAC Tyr	TCC Ser 280	CTT	TAT Tyr	TAC Tyr	AGC Ser	TCT Ser 285	CAG Gln	GAG Glu	ATA Ile	GAC Asp	AAA Lys 290	CCT Pro	TTT Phe		1638
		AGC Ser 295															1686

															GGC Gly	1734
		CTT Leu														1782
		AAC Asn														1830
CCC Pro	GAG Glu	AAT Asn	TCT Ser 360	CCA Pro	GAG Glu	ACA Thr	GAA Glu	GTG Val 365	GCC Ala	CTG Leu	TTT Phe	AGG Arg	ATT Ile 370	AGA Arg	GAC Asp	1878
CGA Arg	GAC Asp	TCT Ser 375	GGA Gly	GAA Glu	AAT Asn	GGA Gly	AAA Lys 380	ATG Met	ATT Ile	TGC Cys	TCA Ser	ATT Ile 385	CAG Gln	GAT Asp	GAT Asp	1926
GTT Val	CCT Pro 390	TTT Phe	AAG Lys	CTA Leu	AAA Lys	CCT Pro 395	TCT Ser	GTT Val	GAG Glu	AAT Asn	TTC Phe 400	TAC Tyr	AGG Arg	CTG Leu	GTA Val	1974
ACA Thr 405	GAA Glu	GGG Gly	GCG Ala	CTG Leu	GAC Asp 410	AGA Arg	GAG Glu	ACC Thr	AGA Arg	GCC Ala 415	GAG Glu	TAC Tyr	AAC Asn	ATC Ile	ACC Thr 420	2022
ATC Ile	ACC Thr	ATC Ile	ACA Thr	GAC Asp 425	TTG Leu	GGG Gly	ACT Thr	CCA Pro	AGG Arg 430	CTG Leu	AAA Lys	ACC Thr	GAG Glu	CAG Gln 435	AGC Ser	2070
ATA Ile	ACC Thr	GTG Val	CTG Leu 440	GTG Val	TCG Ser	GAC Asp	GTC Val	AAT Asn 445	GAC Asp	AAC Asn	GCC Ala	CCC Pro	GCC Ala 450	TTC Phe	ACC Thr	2118
CAA Gln	ACC Thr	TCC Ser 455	TAC Tyr	ACC Thr	CTG Leu	TTC Phe	GTC Val 460	CGC Arg	GAG Glu	AAC Asn	AAC Asn	AGC Ser 465	CCC Pro	GCC Ala	CTG Leu	2166
CAC His	ATC Ile 470	GGC Gly	AGT Ser	GTC Val	AGC Ser	GCC Ala 475	ACA Thr	GAC Asp	AGA Arg	GAC Asp	TCG Ser 480	GGC Gly	ACC Thr	AAC Asn	GCC Ala	2214
CAG Gln 485	GTC Val	ACC Thr	TAC Tyr	TCG Ser	CTG Leu 490	CTG Leu	CCG Pro	CCC Pro	CAG Gln	GAC Asp 495	CCG Pro	CAC His	CTG Leu	CCC Pro	CTA Leu 500	2262
ACC Thr	TCC Ser	CTG Leu	GTC Val	TCC Ser 505	ATT Ile	AAC Asn	ACG Thr	GAC Asp	AAC Asn 510	GGC Gly	CAC His	CTG Leu	TTC Phe	GCT Ala 515	CTC Leu	2310
CAG Gln	TCG Ser	CTG Leu	GAC Asp 520	TAC Tyr	GAG Glu	GCC Ala	CTG Leu	CAG Gln 525	GCT Ala	TTC Phe	GAG Glu	TTC Phe	CGC Arg 530	GTG Val	GGC	2358
GCC Ala	ACA Thr	GAC Asp 535	CGC Arg	GGC Gly	TTC Phe	Pro	GCG Ala 540	CTG Leu	AGC Ser	AGC Ser	GAG Glu	GCG Ala 545	CTG Leu	GTG Val	CGA Arg	2406

					GCC Ala											2454
					GCG Ala 570											2502
					GTG Val											2550
					CTG Leu											2598
					TGG Trp											2646
CTG Leu	CTG Leu 630	AGC Ser	GAG Glu	CGC Arg	GAC Asp	GTG Val 635	GCC Ala	AAG Lys	CAC His	AGG Arg	CTA Leu 640	GTG Val	GTG Val	CTG Leu	GTC Val	2694
					CCT Pro 650											2742
					TTC Phe											2790
GCC Ala	CCG Pro	GCC Ala	CAA Gln 680	GCC Ala	CAG Gln	GCC Ala	GAC Asp	TCG Ser 685	CTT Leu	ACC Thr	GTC Val	TAC Tyr	CTG Leu 690	GTG Val	GTG Val	2838
GCA Ala	TTG Leu	GCC Ala 695	TCG Ser	GTG Val	TCT Ser	TCG Ser	CTC Leu 700	TTC Phe	CTC Leu	TTC Phe	TCG Ser	GTG Val 705	TTC Phe	CTG Leu	TTC Phe	2886
GTG Val	GCA Ala 710	GTG Val	CGG Arg	CTG Leu	TGC Cys	AGG Arg 715	AGG Arg	AGC Ser	AGG Arg	GCG Ala	GCC Ala 720	TCA Ser	GTG Val	GGT Gly	CGC Arg	2934
TGC Cys 725	TCG Ser	GTG Val	CCC Pro	GAG Glu	GGC Gly 730	CCC Pro	TTT Phe	CCA Pro	GGG Gly	CAT His 735	CTG Leu	GTG Val	GAC Asp	GTG Val	AGC Ser 740	2982
GGC Gly	ACC Thr	GGG Gly	ACC Thr	CTT Leu 745	TCC Ser	CAG Gln	AGC Ser	TAC Tyr	CAG Gln 750	TAC Tyr	GAG Glu	GTG Val	TGT Cys	CTG Leu 755	ACG Thr	3030
GGA Gly	GGC Gly	TCT Ser	GAA Glu 760	AGT Ser	AAT Asn	GAT Asp	TTC Phe	AAG Lys 765	TTC Phe	TTG Leu	AAG Lys	CCT Pro	ATA Ile 770	TTC Phe	CCA Pro	3078
AAT Asn	ATT Ile	GTA Val 775	AGC Ser	CAG Gln	GAC Asp	TCT Ser	AGG Arg 780	AGG Arg	AAA Lys	TCA Ser	GAA Glu	TTT Phe 785	CTA Leu	GAA Glu		3123
TAAT	GTAG	GT A	TCTG	TAGO	T TI	CCGA	CCG1	CTG	TTAA	TTT	TGTC	TTCC	TC A	CTTI	TCACC	3183

TTAGTTTTTT	TTAACCCTTT	AGTAATCTTG	AATTCTACTT	TTTTTTAAAT	TTCTACTGTT	3243
GTCTTTAGTA	ATGTTACTCA	TTTCCTTTGT	CTGATTGTTA	GTTTTCAAAT	TATTGTATTA	3303
TTATAAATAT	TTTATATCAG	GAAAGTTCAT	ATTTCTGAAT	AAATTAATAG		3353

(2) INFORMATION FOR SEQ ID NO:110:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 787 amino acids
 - (B) TYPE: amino acid
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: protein
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:110:

Met Glu Pro Ala Gly Glu Arg Phe Pro Glu Gln Arg Gln Val Leu Ile 1 5 10 15

Leu Leu Leu Leu Glu Val Thr Leu Ala Gly Trp Glu Pro Arg Arg
20 25 30

Tyr Ser Val Met Glu Glu Thr Glu Arg Gly Ser Phe Val Ala Asn Leu 35 40 45

Ala Asn Asp Leu Gly Leu Gly Val Gly Glu Leu Ala Glu Arg Gly Ala 50 . 55 60

Arg Val Val Ser Glu Asp Asn Glu Gln Gly Leu Gln Leu Asp Leu Gln 65 70 75 80

Thr Gly Gln Leu Ile Leu Asn Glu Lys Leu Asp Arg Glu Lys Leu Cys 85 90 95

Gly Pro Thr Glu Pro Cys Ile Met His Phe Gln Val Leu Leu Lys Lys
100 105 110

Pro Leu Glu Val Phe Arg Ala Glu Leu Leu Val Thr Asp Ile Asn Asp 115 120 125

His Ser Pro Glu Phe Pro Glu Arg Glu Met Thr Leu Lys Ile Pro Glu 130 135 140

Thr Ser Ser Leu Gly Thr Val Phe Pro Leu Lys Lys Ala Arg Asp Leu 145 150 155 160

Asp Val Gly Ser Asn Asn Val Gln Asn Tyr Asn Ile Ser Pro Asn Ser 165 170 175

His Phe His Val Ser Thr Arg Thr Arg Gly Asp Gly Arg Lys Tyr Pro 180 185 190

Glu Leu Val Leu Asp Thr Glu Leu Asp Arg Glu Glu Gln Ala Glu Leu 195 200 205

Arg Leu Thr Leu Thr Ala Val Asp Gly Gly Ser Pro Pro Arg Ser Gly 210 215 220

Thr Val Gln Ile Leu Ile Leu Val Leu Asp Ala Asn Asp Asn Ala Pro Glu Phe Val Gln Ala Leu Tyr Glu Val Gln Val Pro Glu Asn Ser Pro 245 250 Val Gly Ser Leu Val Val Lys Val Ser Ala Arg Asp Leu Asp Thr Gly Thr Asn Gly Glu Ile Ser Tyr Ser Leu Tyr Tyr Ser Ser Gln Glu Ile Asp Lys Pro Phe Glu Leu Ser Ser Leu Ser Gly Glu Ile Arg Leu Ile 295 Lys Lys Leu Asp Phe Glu Thr Met Ser Ser Tyr Asp Leu Asp Ile Glu Ala Ser Asp Gly Gly Leu Ser Gly Lys Cys Ser Val Ser Val Lys Val Leu Asp Val Asn Asp Asn Phe Pro Glu Leu Ser Ile Ser Ser Leu Thr Ser Pro Ile Pro Glu Asn Ser Pro Glu Thr Glu Val Ala Leu Phe Arg Ile Arg Asp Arg Asp Ser Gly Glu Asn Gly Lys Met Ile Cys Ser Ile Gln Asp Asp Val Pro Phe Lys Leu Lys Pro Ser Val Glu Asn Phe Tyr Arg Leu Val Thr Glu Gly Ala Leu Asp Arg Glu Thr Arg Ala Glu Tyr Asn Ile Thr Ile Thr Ile Thr Asp Leu Gly Thr Pro Arg Leu Lys Thr Glu Gln Ser Ile Thr Val Leu Val Ser Asp Val Asn Asp Asn Ala Pro Ala Phe Thr Gln Thr Ser Tyr Thr Leu Phe Val Arg Glu Asn Asn 455 Ser Pro Ala Leu His Ile Gly Ser Val Ser Ala Thr Asp Arg Asp Ser Gly Thr Asn Ala Gln Val Thr Tyr Ser Leu Leu Pro Pro Gln Asp Pro 485 490 His Leu Pro Leu Thr Ser Leu Val Ser Ile Asn Thr Asp Asn Gly His 500 Leu Phe Ala Leu Gln Ser Leu Asp Tyr Glu Ala Leu Gln Ala Phe Glu Phe Arg Val Gly Ala Thr Asp Arg Gly Phe Pro Ala Leu Ser Ser Glu Ala Leu Val Arg Val Leu Val Leu Asp Ala Asn Asp Asn Ser Pro Phe 545 550 555 560

Val Leu Tyr Pro Leu Gln Asn Gly Ser Ala Pro Cys Thr Glu Leu Val 565 570 575

Pro Arg Ala Ala Glu Pro Gly Tyr Leu Val Thr Lys Val Val Ala Val 580 585 590

Asp Gly Asp Ser Gly Gln Asn Ala Trp Leu Ser Tyr Gln Leu Lys 595 600 605

Ala Thr Glu Pro Gly Leu Phe Gly Val Trp Ala His Asn Gly Glu Val 610 620

Arg Thr Ala Arg Leu Leu Ser Glu Arg Asp Val Ala Lys His Arg Leu 625 630 635 640

Val Val Leu Val Lys Asp Asn Gly Glu Pro Pro Arg Ser Ala Thr Ala 645 650 655

Thr Leu Gln Val Leu Leu Val Asp Gly Phe Ser Gln Pro Tyr Leu Pro 660 665 670

Leu Pro Glu Ala Ala Pro Ala Gln Ala Gln Ala Asp Ser Leu Thr Val 675 680 685

Tyr Leu Val Val Ala Leu Ala Ser Val Ser Ser Leu Phe Leu Phe Ser 690 695 700

Val Phe Leu Phe Val Ala Val Arg Leu Cys Arg Arg Ser Arg Ala Ala 705 710 715 720

Ser Val Gly Arg Cys Ser Val Pro Glu Gly Pro Phe Pro Gly His Leu
725 730 735

Val Asp Val Ser Gly Thr Gly Thr Leu Ser Gln Ser Tyr Gln Tyr Glu 740 745 750

Val Cys Leu Thr Gly Gly Ser Glu Ser Asn Asp Phe Lys Phe Leu Lys 755 760 765

Pro Ile Phe Pro Asn Ile Val Ser Gln Asp Ser Arg Arg Lys Ser Glu
770 780

Phe Leu Glu 785

(2) INFORMATION FOR SEQ ID NO:111:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 3033 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA

(ix) FEATURE:
(A) NAME/KEY: CDS
(B) LOCATION: 138..2528

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11	(xi)	SEQUENCE	DESCRIPTION:	SEQ	ID	NO:111
---	------	----------	--------------	-----	----	--------

GTGA	TTG	GAC (GTGT	TTTT	GT G	ACTA	TTTG	G GA	AGAA	GACA	CCT	TCCT	AAT (CAGA	TTTACT	60
CCAA	TAT	CTT (CCCG	GACC	CT C	ATGA	GTGG	A TT	GCAA	TTGA	CTT	GAAG.	AAG (CAGC	ACCCTC	120
AGGA	.CTG/	AAT (CTGA											CAG (Gln (170
AGG (CAA Gln	GTC Val	TTT Phe 15	TTT Phe	CTT Leu	ACT	ATA Ile	TTG Leu 20	TCG Ser	TTA Leu	TTG Leu	TGG Trp	AAG Lys 25	TCT Ser	AGC Ser	218
TCT (GAG Glu	GCC Ala 30	ATT Ile	AGA Arg	TAT Tyr	TCC Ser	ATG Met 35	CCA Pro	GAA Glu	GAA Glu	ACA Thr	GAG Glu 40	AGT Ser	GGC	TAT Tyr	266
ATG (314
TCC Ser S	TCT Ser	AGA Arg	GGA Gly	GCT Ala	CAA Gln 65	ATC Ile	CAT His	TAC Tyr	AAA Lys	GGA Gly 70	AAC Asn	AAA Lys	GAA Glu	CTT Leu	TTG Leu 75	362
CAG (CTG Leu	GAT Asp	GCA Ala	GAG Glu 80	ACT Thr	GGG Gly	AAT Asn	TTG Leu	TTC Phe 85	TTA Leu	AAG Lys	GAA Glu	AAA Lys	CTA Leu 90	GAC Asp	410
AGA (GAA Glu	CTG Leu	CTG Leu 95	TGT Cys	GGA Gly	GAG Glu	ACA Thr	GAA Glu 100	CCC Pro	TGT Cys	GTG Val	CTG Leu	AAC Asn 105	TTC Phe	CAG Gln	458
ATC I	ATA Ile	CTG Leu 110	GAA Glu	AAC Asn	CCT Pro	ATG Met	CAG Gln 115	TTC Phe	TTC Phe	CAA Gln	ACT Thr	GAA Glu 120	CTG Leu	CAG Gln	CTC Leu	506
ACA C	GAT Asp 125	ATA Ile	AAC Asn	GAC Asp	CAT His	TCT Ser 130	CCA Pro	GAG Glu	TTC Phe	CCC Pro	AAC Asn 135	AAG Lys	AAA Lys	ATG Met	CTT Leu	554
CTA I Leu 1 140	ACA Thr	ATT Ile	CCT Pro	GAG Glu	AGT Ser 145	GCC Ala	CAT His	CCA Pro	GGG Gly	ACT Thr 150	GTG Val	TTT Phe	CCT Pro	CTG Leu	AAG Lys 155	602
GCA (GCT Ala	CGG Arg	GAC Asp	TCT Ser 160	GAC Asp	ATA Ile	GCG	AGC Ser	AAC Asn 165	GCT Ala	GTT Val	CAG Gln	AAC Asn	TAC Tyr 170	ACA Thr	650
GTC P	AAT Asn	CCC Pro	AAC Asn 175	CTC Leu	CAT His	TTC Phe	CAC His	GTC Val 180	GTT Val	ACT Thr	CAC His	AGT Ser	CGC Arg 185	ACA Thr	GAT Asp	698

					GAG Glu											746
					ACT Thr											794
					ACC Thr 225											842
					CAG Gln											890
					CTC Leu											938
					GTA Val											986
					CAA Gln											1034
					GAG Glu 305											1082
					ACA Thr											1130
					TTG Leu											1178
					GTC Val											1226
GTT Val	GCT Ala 365	GTT Val	TTT Phe	AGT Ser	GTT Val	TCT Ser 370	GAT Asp	TCT Ser	GAT Asp	TCG Ser	GGG Gly 375	GAC Asp	AAT Asn	GGA Gly	AGG Arg	1274
					CCG Pro 385											1322
					ACG Thr											1370
AAC Asn	AGA Arg	GCT Ala	GAG Glu 415	TAC Tyr	AAC Asn	ATC Ile	ACC Thr	ATC Ile 420	ACG Thr	GTC Val	TCA Ser	GAT Asp	CTG Leu 425	GGC Gly	ACA Thr	1418

CCC Pro	AGG Arg	Leu 430	Thr	ACC Thr	CAG Gln	CAC His	ACC Thr 435	Ile	ACA Thr	GTG Val	CAA Gln	GTG Val 440	Ser	GAC	ATC Ile	•	1466
AAC Asn	GAC Asp 445	Asn	GCC Ala	CCT Pro	GCC Ala	TTC Phe 450	ACC Thr	CAA Gln	ACC Thr	TCC ,Ser	TAC Tyr 455	Thr	ATG Met	TTT Phe	GTC Val		1514
CAC His 460	Glu	AAC	AAC Asn	AGC Ser	Pro 465	GCC Ala	CTG Leu	CAC His	ATA Ile	GGC Gly 470	ACC Thr	ATC	AGT Ser	GCC Ala	ACA Thr 475		1562
GAC Asp	TCA Ser	GAC	TCA Ser	GGC Gly 480	Ser	AAT Asn	GCC Ala	CAC His	ATC Ile 485	ACC Thr	TAC Tyr	TCG Ser	CTG Leu	CTG Leu 490	CCG Pro		1610
Pro	Asp	Asp	Pro 495	Gln	Leu	GCC Ala	Leu	Asp 500	Ser	Leu	Ile	Ser	Ile 505	Asn	Val		1658
GAC Asp	AAT Asn	GGG Gly 510	CAG Gln	CTG Leu	TTC Phe	GCG Ala	CTC Leu 515	AGA Arg	GCT Ala	CTA Leu	GAC Asp	TAT Tyr 520	GAG Glu	GCA Ala	CTG Leu		1706
CAG Gln	TCC Ser 525	TTC Phe	GAG Glu	TTC Phe	TAC Tyr	GTG Val 530	GCG	GCT Ala	ACA Thr	GAT Asp	GGA Gly 535	GGC Gly	TCA Ser	CCC Pro	GCG Ala		1754
CTC Leu 540	AGC Ser	AGC Ser	CAG Gln	ACT Thr	CTG Leu 545	GTG Val	CGG Arg	ATG Met	GTG Val	GTG Val 550	CTG Leu	GAT Asp	GAC Asp	AAT Asn	GAC Asp 555		1802
Asn	Ala	Pro	Phe	Val 560	Leu	TAC Tyr	Pro	Leu	Gln 565	Asn	Ala	Ser	Ala	Pro 570	Cys		1850
ACT Thr	GAG Glu	CTA Leu	CTG Leu 575	CCT Pro	AGG Arg	GCA Ala	GCA Ala	GAG Glu 580	CCC Pro	GGC Gly	TAC Tyr	CTG Leu	ATC Ile 585	ACC Thr	AAA Lys		1898
GTG Val	GTG Val	GCT Ala 590	GTG Val	GAT Asp	CGC Arg	GAC Asp	TCT Ser 595	GGA Gly	CAG Gln	AAT Asn	GCT Ala	TGG Trp 600	CTG Leu	TCG Ser	TTC Phe		1946
CAG Gln	CTA Leu 605	CTT Leu	AAA Lys	GCT Ala	ACA Thr	GAG Glu 610	CCA Pro	GGG Gly	CTG Leu	TTC Phe	AGT Ser 615	GTA Val	TGG Trp	GCA Ala	CAC His		1994
AAT Asn 620	GGT Gly	GAA Glu	GTG Val	CGC Arg	ACC Thr 625	ACT Thr	AGG Arg	CTG Leu	CTG Leu	AGT Ser 630	GAG Glu	CGA Arg	GAT Asp	GCT Ala	CAG Gln 635		2042
AAG Lys	nls	ràs	Leu	Leu 640	Leu	Leu	Val	Lys	Asp 645	Asn	Gly	Asp	Pro	Leu 650	Arg		2090
TCT Ser	GCC Ala	AAT Asn	GTC Val 655	ACT Thr	CTT Leu	CAC (Val	CTA Leu 660	GTG Val	GTG Val	GAT Asp	Gly	TTC Phe 665	TCG Ser	CAG Gln		2138

CCT Pro	TAC Tyr	CTA Leu 670	CCA Pro	TTG Leu	GCT Ala	GAG Glu	GTG Val 675	GCA Ala	CAG Gln	GAT Asp	TCC Ser	ATG Met 680	CAA Gln	GAT Asp	AAT Asn	2186
TAC Tyr	GAC Asp 685	GTT Val	CTC Leu	ACA Thr	CTG Leu	TAC Tyr 690	CTA Leu	GTC Val	ATT Ile	GCC Ala	TTG Leu 695	GCA Ala	TCT Ser	GTA Val	TCT Ser	2234
TCT Ser 700	CTC Leu	TTC Phe	CTC Leu	TTG Leu	TCT Ser 705	GTA Val	GTG Val	CTG Leu	TTT Phe	GTG Val 710	GGG Gly	GTG Val	AGG Arg	CTG Leu	TGC Cys 715	2282
AGG Arg	AGG Arg	GCC Ala	AGG Arg	GAG Glu 720	GCC Ala	TCC Ser	TTG Leu	GGT Gly	GAC Asp 725	TAC Tyr	TCT Ser	GTG Val	CCT Pro	GAG Glu 730	GGA Gly	2330
CAC His	TTT Phe	CCT Pro	AGC Ser 735	CAC His	TTG Leu	GTG Val	GAT Asp	GTC Val 740	AGC Ser	GGT Gly	GCC Ala	GGG Gly	ACC Thr 745	CTG Leu	TCC Ser	2378
CAG Gln	AGT Ser	TAT Tyr 750	CAA Gln	TAT Tyr	GAG Glu	GTG Val	TGT Cys 755	CTT Leu	AAT Asn	GGA Gly	GGT Gly	ACT Thr 760	AGA Arg	ACA Thr	AAT Asn	2426
GAG Glu	TTT Phe 765	AAC Asn	TTT Phe	CTT Leu	AAA Lys	CCA Pro 770	TTG Leu	TTT Phe	CCT Pro	ATC Ile	CTT Leu 775	CCG Pro	ACC Thr	CAG Gln	GCT Ala	2474
GCT Ala 780	GCT Ala	GCT Ala	GAA Glu	GAA Glu	AGA Arg 785	GAA Glu	AAC Asn	GCT Ala	GTT Val	GTG Val 790	CAC His	AAT Asn	AGC Ser	GTT Val	GGA Gly 795	2522
TTC Phe	TAT Tyr	TAGA	GCAC	TG A	TTTT	'GAAG	T GG	TGGT	TACC	TCA	\TTTT	TCC	TTAA	CTAT	CC	2578
CTGA	TGTA	GA A	TGGT	GTAG	T GC	CGTG	AATC	AAC	TCCT	GAG	ATAT	ATGT	TC A	TTTT	ATCCT	2638
TTGT	TTTG.	AA T	CAAA	CTAT	T CA	GATG	TGAT	CCT	ACTC	TAG	AGAA	TTTG	GT I	CTAC	TCCAT	2698
TGTG'	TTTG	TT T	AGAT	TTCT	A CG	CCAT	ACCA	GTG	CATG	CTG	GGTT	GTTT	TT T	TTTT	TACAA	2758
TAT'	TATA	AC T	TTGC	TTTG	G AG	GGGA	ACTC	ATA	TTCG	CTG	TAAC	GAAT	TG G	AACC	ACTTT	2818
CATT	GTTA	GA G	ATGC	CTTG	C TT	TGTT	GTGT	TAT	TTCA	GAC	AGGG	TCTT	AA A	TTGT	AGCCC	2878
rggg:	TGAC	CT G	AAAT	GACT.	A TG	TACA	GACT	GAC	TTTG.	AAT	TTGT	GGCA	GT C	CATC	TGCCT	2938
CTGT	TGTC	CT A	TGTT	GGGA	T TG	TGAG	CATG	CAT	GAGT.	AGG	CTCA	GCTG	TG G	TGAG	CGACC	2998
TAA!	TAAA	AA T	CAAA	TACT	A AA	AAAA	AAAA	AAA	AA							3033

(2) INFORMATION FOR SEQ ID NO:112:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 797 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:112:

Tyr Ser Met Pro Glu Glu Thr Glu Ser Gly Tyr Met Val Ala Asn Leu
35 40 45

Ala Lys Asp Leu Gly Ile Arg Val Gly Glu Leu Ser Ser Arg Gly Ala 50 60

Gln Ile His Tyr Lys Gly Asn Lys Glu Leu Leu Gln Leu Asp Ala Glu 65 70 75 80

Thr Gly Asn Leu Phe Leu Lys Glu Lys Leu Asp Arg Glu Leu Leu Cys 85 90 95

Gly Glu Thr Glu Pro Cys Val Leu Asn Phe Gln Ile Ile Leu Glu Asn 100 105 110

Pro Met Gln Phe Phe Gln Thr Glu Leu Gln Leu Thr Asp Ile Asn Asp 115 120 125

His Ser Pro Glu Phe Pro Asn Lys Lys Met Leu Leu Thr Ile Pro Glu 130 135 140

Ser Ala His Pro Gly Thr Val Phe Pro Leu Lys Ala Ala Arg Asp Ser 145 150 155 160

Asp Ile Gly Ser Asn Ala Val Gln Asn Tyr Thr Val Asn Pro Asn Leu 165 170 175

His Phe His Val Val Thr His Ser Arg Thr Asp Gly Arg Lys Tyr Pro 180 185 190

Glu Leu Val Leu Asp Arg Ala Leu Asp Arg Glu Glu Gln Pro Glu Leu 195 200 205

Thr Leu Ile Leu Thr Ala Leu Asp Gly Gly Ala Pro Ser Arg Ser Gly 210 215 220

Thr Thr Thr Val His Ile Glu Val Val Asp Ile Asn Asp Asn Ser Pro 225 230 240

Gln Phe Val Gln Ser Leu Tyr Lys Val Gln Val Pro Glu Asn Asn Pro 245 250 255

Leu Asn Ala Phe Val Val Thr Val Ser Ala Thr Asp Leu Asp Ala Gly 260 265 270

Val Tyr Gly Asn Val Thr Tyr Ser Leu Phe Gln Gly Tyr Gly Val Phe 275 280 285

Gln Pro Phe Val Ile Asp Glu Ile Thr Gly Glu Ile His Leu Ser Lys 290 295 300

Glu Leu Asp Phe Glu Glu Ile Ser Asn His Asn Ile Glu Ile Ala Ala Thr Asp Gly Gly Leu Ser Gly Lys Cys Thr Val Ala Val Gln Val Leu Asp Val Asn Asp Asn Ala Pro Glu Leu Thr Ile Arg Lys Leu Thr Val Leu Val Pro Glu Asn Ser Ala Glu Thr Val Val Ala Val Phe Ser Val Ser Asp Ser Asp Ser Gly Asp Asn Gly Arg Met Val Cys Ser Ile Pro Asn Asn Ile Pro Phe Leu Leu Lys Pro Thr Phe Glu Asn Tyr Tyr Thr Leu Val Thr Glu Gly Pro Leu Asp Arg Glu Asn Arg Ala Glu Tyr Asn Ile Thr Ile Thr Val Ser Asp Leu Gly Thr Pro Arg Leu Thr Thr Gln His Thr Ile Thr Val Gln Val Ser Asp Ile Asn Asp Asn Ala Pro Ala Phe Thr Gln Thr Ser Tyr Thr Met Phe Val His Glu Asn Asn Ser Pro Ala Leu His Ile Gly Thr Ile Ser Ala Thr Asp Ser Asp Ser Gly Ser Asn Ala His Ile Thr Tyr Ser Leu Leu Pro Pro Asp Asp Pro Gln Leu Ala Leu Asp Ser Leu Ile Ser Ile Asn Val Asp Asn Gly Gln Leu 505 Phe Ala Leu Arg Ala Leu Asp Tyr Glu Ala Leu Gln Ser Phe Glu Phe Tyr Val Gly Ala Thr Asp Gly Gly Ser Pro Ala Leu Ser Ser Gln Thr Leu Val Arg Met Val Val Leu Asp Asp Asn Asp Asn Ala Pro Phe Val Leu Tyr Pro Leu Gln Asn Ala Ser Ala Pro Cys Thr Glu Leu Leu Pro 570 Arg Ala Ala Glu Pro Gly Tyr Leu Ile Thr Lys Val Val Ala Val Asp 585 Arg Asp Ser Gly Gln Asn Ala Trp Leu Ser Phe Gln Leu Leu Lys Ala Thr Glu Pro Gly Leu Phe Ser Val Trp Ala His Asn Gly Glu Val Arg

Thr 625	Thr	Arg	Leu	Leu	Ser 630	Glu	Arg	Asp	Ala	Gln 635	Lys	His	Lys	Leu	Leu 640
Leu	Leu	Val	Lys	Asp 645	Asn	Gly	Asp	Pro	Leu 650	Arg	Ser	Ala	Asn	Val 655	Thr
Leu	His	Val	Leu 660	Val	Val	Asp	Gly	Phe 665	Ser	Gln	Pro	Tyr	Leu 670	Pro	Leu
Ala	Glu	Val 675	Ala	Gln	Asp	Ser	Met 680	Gln	Asp	Asn	Tyr	Asp 685	Val	Leu	Thr
Leu	Tyr 690	Leu	Val	Ile	Ala	Leu 695	Ala	Ser	Val	Ser	Ser 700	Leu	Phe	Leu	Leu
Ser 705	Val	Val	Leu	Phe	Val 710	Gly	Val	Arg	Leu	Cys 715	Arg	Arg	Ala	Arg	Glu 720
Ala	Ser	Leu	Gly	Asp 725	Tyr	Ser	Val	Pro	Glu 730	Gly	His	Phe	Pro	Ser 735	His
Leu	Val	Asp	Val 740	Ser	Gly	Ala	Gly	Thr 745	Leu	Ser	Gln	Ser	Tyr 750	Gln	Tyr
Glu	Val	Cys 755	Leu	Asn	Gly	Gly	Thr 760	Arg	Thr	Asn	Glu	Phe 765	Asn	Phe	Leu
Lys	Pro 770	Leu	Phe	Pro	Ile	Leu 775	Pro	Thr	Gln	Ala	Ala 780	Ala	Ala	Glu	Glu
Arg 785	Glu	Asn	Ala	Val	Val 790	His	Asn	Ser	Val	Gly 795	Phe	Tyr			

- (2) INFORMATION FOR SEQ ID NO:113:
 - (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 2347 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear
 - (ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:113:

AAAACACGGG	GGAAATGACA	GTAGCAAAGA	ATCTGGACTA	TGAAGAATGC	TCATTGTATG	60
AAATGGAAAT	ACAGGCTGAA	GATGTGGGGG	CGCTTCTGGG	GAGGAGCAAA	GTGGTAATTA	120
TGGTAGAAGA	TGTAAATGAC	AATCGGCCAG	AAGTGACCAT	TACATCCTTG	TTTAACCCGG	180
TATTGGAAAA	TTCTCTTCCC	GGGACAGTAA	TTGCCTTCTT	GAATGTGCAT	GACCGAGACT	240
CTGGAAAGAA	CGGCCAAGTT	GTCTGTTACA	CGCATGATAA	CTTACCTTTT	AAATTAGAAA	300
AGTCAATAGA	TAATTATTAT	AGATTGGTGA	CATGGAAATA	TTTGGACCGA	GAAAAAGTCT	360
CCATCTACAA	TATCACAGTG	ATAGCCTCAG	ATCTAGGAGC	CCACTCTGTC	ACTGAAACTT	420

ACATTGCCCT	GATTGTGGCA	GACACTAATG	ACAACCCTCC	TCGTTTTCCT	CACACCTCCT	480
ACACAGCCTA	TATTCCAGAG	AACAACCTGA	GGGGCGCCTC	CATCTTCTCA	CTGACTGCAC	540
ATGATCCTGA	CAGTCAGGAA	AATGCACAGG	TCACTTACTC	TGTGTCTGAG	GACACCATAC	600
AGGGAGTGCC	TTTGTCCTCT	TATATCTCCA	TCAACTCAGA	TACTGGTGTC	CTGTATGCAC	660
TGCACTCTTT	TGACTTCGAG	AAGATACAAG	ACTTGCAGCT	ACTGGTTGTT	GCCACTGACA	720
GTGGAAGCCC	ACCTCTCAGC	AGCAATGTGT	CATTGAGCTT	GTTTGTGTTG	GACCAGAACG	780
ACAACGCACC	TGAGATTCTA	TATCCTAGCT	TCCCCACAGA	TGGCTCCACT	GGTGTGGAAC	840
TAGCACCCCG	CTCTGCAGAG	CCTGGATACC	TAGTGACCAA	AGTGGTGGCA	GTGGACAAAG	900
ACTCAGGACA	GAATGCTTGG	CTGTCCTACC	GTCTGCTGAA	GGCCAGCGAA	CCTGGGCTCT	960
TCTCTGTAGG	ACTTCACACG	GGTGAGGTGC	GTACAGCGAG	GGCCCTGCTG	GACAGAGATG	1020
CTCTCAAACA	GAATCTGGTG	ATGGCCGTGC	AGGACCATGG	CCAACCCCCT	CTCTCGGCCA	1080
CTGTAACTCT	CACTGTGGCA	GTGGCTAACA	GCATCCCTGA	GGTGTTGGCT	GACTTGAGCA	1140
GCATTAGGAC	CCCTGGGGTA	CCAGAGGATT	CTGATATCAC	GCTCCACCTG	GTGGTGGCAG	1200
TGGCTGTGGT	CTCCTGTGTC	TTCCTTGTCT	TTGTCATTGT	CCTCCTAGCT	CTCAGGCTTC	1260
AGCGCTGGCA	GAAGTCTCGC	CAGCTCCAGG	GCTCCAAAGG	TGGATTGGCT	CCTGCACCTC	1320
CATCACATTT	TGTGGGCATC	GACGGGGTAC	AGGCTTTTCT	ACAAACCTAT	TCTCATGAAG	1380
TCTCGCTCAC	TTCAGGCTCC	CAGACAAGCC	ACATTATCTT	TCCTCAGCCC	AACTATGCAG	1440
ACATGCTCAT	TAACCAAGAA	GGCTGTGAGA	AAAATGATTC	CTTATTAACA	TCCATAGATT	1500
TTCATGAGAG	TAACCGTGAA	GATGCTTGCG	CCCCGCAAGC	CCCGCCCAAC	ACTGACTGGC	1560
GTTTCTCTCA	AGCCCAGAGA	CCCGGCACGA	GCGGATCCCA	AAATGGGGAT	GAAACCGGCA	1620
CCTGGCCCAA	CAACCAGTTC	GATACAGAGA	TGCTGCAAGC	CATGATCTTG	GCCTCTGCCA	1680
GTGAAGCCGC	TGATGGGAGC	TCCACTCTGG	GAGGGGGCAC	TGGCACTATG	GGTTTGAGCG	1740
CTCGATATGG	ACCCCAGTTT	ACCCTGCAGC	ACGTGCCTGA	CTACCGCCAG	AACGTGTACA	1800
TCCCTGGCAG	CAATGCCACA	CTGACCAACG	CAGCTGGCAA	ACGAGATGGC	AAGGCTCCGG	1860
CAGGCGGCAA	TGGCAACAAC	AACAAGTCGG	GCAAGAAAGA	GAAGAAGTAA	TATGGAGGCC	1920
AGGCCTTGAG	CCACAGGGCA	GCCTCCCTCC	CCAGCCAGTC	CAGCTTGTCC	TTACTTGTAC	1980
CCAGGCCTCA	GAATTTCAGG	GCTCACCCCA	GGATTCTGGT	AGGAGCCACA	GCCAGGCCAT	2040
GCTCCCCGTT	GGGAAACAGA	AACAAGTGCC	CAAGCCAACA	CCCCCTCTTT	GTACCCTAGG	2100
GGGGTTGAAT	ATGCAAAGAG	AGTTCTGCTG	GGACCCCCTA	TCCAATCAGT	GATTGTACCC	2160
ACATAGGTAG	CAGGGTTAGT	GTGGATACAC	ACACACACAC	ACACACACAC	ACACACACAA	2220
CCCTTGTCCT	CCGCAGTGCC	TGCCACTTTC	TGGGACTTTC	TCATCCCCCT	ACGCCCTTCC	2280

TTT	CATCO	CTCT	CCCF	CCCA	GA C	CACAG	CTGC	T GG	AGAA	TAAA	TTI	'GGGG	ATG	CTGA	TGCT.	AA 2340
AAA	AAAA	\														2347
(2)	INF	FORMA	MOITA	FOR	SEÇ] ID	NO: 1	14:								
	(<u>i</u>	(A) I B) I C) S	ENGT YPE: TRAN	H: 2 nuc DEDN	CTER 1972 :leic ESS: lin	base aci sin	pai d	.rs							
	(ii	.) MC	LECU	LE T	YPE:	CDN	A									
	(ix	•	A) N	AME/		CDS 2										
	(xi) SE	QUEN	CE D	ESCR	IPTI	on:	SEQ	ID N	0:11	4:					
A G	AG G lu A 1	CT G la A	CT C la H	AC C is H	AC C is L 5	TG G eu V	TC C al L	TC A eu T	hr A	CC To	CG G er A	AT G	GC G ly G	ly L	AG ys 15	46
CCG Pro	CCT Pro	CGC Arg	TCT Ser	AGC Ser 20	Thr	GTG Val	CGC Arg	ATC Ile	CAC His 25	GTG Val	ACA Thr	GTG Val	TTG Leu	GAT Asp 30	ACA Thr	94
AAT Asn	GAC Asp	AAT Asn	GCC Ala 35	CCG Pro	GTT Val	TTT Phe	CCT Pro	CAC His 40	CCG Pro	ATT Ile	TAC Tyr	CGA Arg	GTG Val 45	AAA Lys	GTC Val	142
CTT Leu	GAG Glu	AAC Asn 50	ATG Met	CCC	CCA Pro	GGC Gly	ACG Thr 55	CGG Arg	CTG Leu	CTT Leu	ACT Thr	GTA Val 60	ACA Thr	GCC Ala	AGC Ser	190
GAC Asp	CCG Pro 65	GAT Asp	GAG Glu	GGA Gly	ATC Ile	AAC Asn 70	GGA Gly	AAA Lys	GTG Val	GCA Ala	TAC Tyr 75	AAA Lys	TTC Phe	CGG Arg	AAA Lys	238
ATT Ile 80	AAT Asn	GAA Glu	AAA Lys	CAA Gln	ACT Thr 85	CCG Pro	TTA Leu	TTC Phe	CAG Gln	CTT Leu 90	AAT Asn	GAA Glu	AAT Asn	ACT Thr	GGG Gly 95	286
GAA Glu	ATA Ile	TCA Ser	ATA Ile	GCA Ala 100	AAA Lys	AGT Ser	CTA Leu	GAT Asp	TAT Tyr 105	GAA Glu	GAA Glu	TGT Cys	TCA Ser	TTT Phe 110	TAT Tyr	334
GAA Glu	ATG Met	GAA Glu	ATA Ile 115	CAA Gln	GCC Ala	GAA Glu	GAT Asp	GTG Val 120	GGG Gly	GCA Ala	CTT Leu	CTG Leu	GGG Gly 125	AGG Arg	ACC Thr	382
AAA Lys	TTG Leu	CTC Leu 130	ATT Ile	TCT Ser	GTG Val	GAA Glu	GAT Asp 135	GTA Val	AAT Asn	GAC Asp	AAT Asn	AGA Arg 140	CCA Pro	GAA Glu	GTG Val	430
ATC Ile	ATT Ile 145	ACG Thr	TCT Ser	TTG Leu	TTT Phe	AGC Ser 150	CCA Pro	GTG Val	TTA Leu	GAA Glu	AAT Asn 155	TCT Ser	CTT Leu	CCC Pro	GGG Gly	478

ACA Thr 160	Val	ATT Ile	GCC Ala	TTC Phe	TTG Leu 165	AGT Ser	GTG Val	CAT	GAC Asp	CAA Gln 170	GAC Asp	TCT Ser	GGA Gly	AAG Lys	AAT Asn 175	526
GGT Gly	CAA Gln	GTT Val	GTC Val	TGT Cys 180	Tyr	ACA Thr	CGT	GAT Asp	AAT Asn 185	TTA Leu	CCT	TTT Phe	AAA Lys	TTA Leu 190		574
AAG Lys	TCA Ser	ATA Ile	GGT Gly 195	AAT Asn	TAT Tyr	TAT Tyr	AGA Arg	TTA Leu 200	GTG Val	ACA Thr	AGG Arg	AAA Lys	TAT Tyr 205	TTG Leu	GAC Asp	622
CGA Arg	GAA Glu	AAT Asn 210	Val	TCT Ser	ATC Ile	TAC Tyr	AAT Asn 215	ATC Ile	ACA Thr	GTG Val	ATG Met	GCC Ala 220	TCA Ser	GAT Asp	CTA Leu	670
GGA Gly	ACA Thr 225	CCA Pro	CCT	CTG Leu	TCC Ser	ACT Thr 230	GAA Glu	ACT Thr	CAA Gln	ATC Ile	GCT Ala 235	CTG Leu	CAC His	GTG Val	GCA Ala	718
GAC Asp 240	Ile	AAC Asn	GAC Asp	AAC Asn	CCT Pro 245	CCT Pro	ACT Thr	TTC Phe	CCT Pro	CAT His 250	GCC Ala	TCC Ser	TAC Tyr	TCA Ser	GCG Ala 255	766
TAT Tyr	ATC Ile	CTA Leu	GAG Glu	AAC Asn 260	AAC Asn	CTG Leu	AGA Arg	GGA Gly	GCC Ala 265	TCC Ser	ATC Ile	TTT Phe	TCC Ser	TTG Leu 270	ACT Thr	814
GCA Ala	CAC His	GAC Asp	CCC Pro 275	GAC Asp	AGC Ser	CAG Gln	GAG Glu	AAT Asn 280	GCC Ala	CAG Gln	GTC Val	ACT Thr	TAC Tyr 285	TCT Ser	GTG Val	862
ACC Thr	GAG Glu	GAC Asp 290	ACG Thr	CTG Leu	CAG Gln	GGG Gly	GCG Ala 295	CCC Pro	CTG Leu	TCC Ser	TCG Ser	TAT Tyr 300	ATC Ile	TCC Ser	ATC Ile	910
AAC Asn	TCT Ser 305	GAC Asp	ACC Thr	GGT Gly	GTC Val	CTG Leu 310	TAT Tyr	GCG Ala	CTG Leu	CAA Gln	TCT Ser 315	TTC Phe	GAC Asp	TAT Tyr	GAG Glu	958
CAG Gln 320	ATC Ile	CGA Arg	GAC Asp	CTG Leu	CAG Gln 325	CTA Leu	CTG Leu	GTA Val	ACA Thr	GCC Ala 330	AGC Ser	GAC Asp	AGC Ser	GGG Gly	GAC Asp 335	1006
CCG Pro	CCC Pro	CTC Leu	AGC Ser	AGC Ser 340	AAC Asn	ATG Met	TCA Ser	CTG Leu	AGC Ser 345	CTG Leu	TTC Phe	GTG Val	CTG Leu	GAC Asp 350	CAG Gln .	1054
AAT Asn	GAC Asp	AAC Asn	GCG Ala 355	CCC Pro	GAG Glu	ATC Ile	CTG Leu	TAC Tyr 360	CCC Pro	GCC Ala	CTC Leu	CCC Pro	ACA Thr 365	GAC Asp	GGT Gly	1102
TCC Ser	ACT Thr	GGC Gly 370	GTG Val	GAG Glu	CTG Leu	GCG Ala	CCC Pro 375	CGC Arg	TCC Ser	GCA Ala	GAG Glu	CGT Arg 380	GGC Gly	TAC Tyr	CTG Leu	1150
GTG Val	ACC Thr 385	AAG Lys	GTG Val	GTG Val	GCG Ala	GTG Val 390	GAC Asp	AGA Arg	GAC Asp	TCG Ser	GGC Gly 395	CAG Gln	AAC Asn	GCC Ala	TGG Trp	1198

			CGC Arg													1246
			ACG Thr													1294
			AAG Lys 435													1342
			TCC Ser													1390
			GTC Val													1438
			TCG Ser													1486
			GTC Val													1534
			TGG Trp 515													1582
			GTG Val													1630
			CAG Gln													1678
			CAC His													1726
			GAG Glu													1774
															TTA Leu	1822
			TTG Leu						TGA	ATTT:	TAT :	rtgg(CATAI	AA		1869
TTA	rgtt:	TTG A	LAAA	ACAT	TG TO	GAAG!	ATAG	r TG?	\AAA?	TAAT	TTT	PAAGO	TG T	TATC	ACAGAG	1929
TTT	rggg:	TTT A	ATTT	CGT	G TO	TTAC	CAA	A AA	ATTG!	ACT	CTA	ATAG	CA 1	ragg:	TATTG	1989
TTTC	CATT	rgc 1	TTTI	AAAC	SA C	TGG	AAAA	AT1	CTTO	CCAC	CAT	(ATT	AAC (CTTC	CAGTAT	2049

TTTATTCCTA	TTATCACTCA	TTCACTTAAG	AAGTAGCTAC	CCGTCCATAC	TGGTAATTTT	2109
GCTATTGTTT	GTTTGTGTGT	GTGTGTGT	GTGTGTGT	GTGTGTGTAT	CCCAAACTAG	2169
AACTTCAGAA	AATTATCAAG	AAGTCTAAAG	CCTTGTTATT	AGCTTAGCAA	AAGTAAAATA	2229
TATCTCAGAA	TTTTTAGGGT	TATGTTTAGC	ATTTGAACCT	GTAACTAGGC	TCTTGTATAT	2289
TTCTTCACTT	TAAACCTCTT	TTCTGAGCCC	TGTTTCTGTA	CCAGTGCCCT	TCAAAACTTT	2349
AATACTTCTT	ACCATCCTTC	AAAACATGAA	CAAACTTTAA	AGATGGATCT	TGGTGGGAGA	2409
TGAGACTGGT	TACTAAATAT	TAAGTATGTG	AGTCAGTGGT	CACCTGGGCT	CCATCCCCAT	2469
GGAGACATGA	AATCTAAAGC	CTAGAATGTC	CATTGCTCCC	CCAAACAAAA	AACAAAAGCA	2529
AAAACATTAG	ATCTGAATTA	AAATGTAATT	TTAAACTGTT	GAAAGTGACT	TTTGTAAAAT	2589
ATGTAAGAAC	ATATTTCAAT	ACAATTCCAA	TTAGCTGTTT	CGGTTGTGCA	TTGATGTGAA	2649
GTGGTGAGAA	TGTTGATATT	AAGAACCAAT	GTTTCAGGTA	CACAAGTTCT	AAATAAGCTG	2709
ATCAATTCAA	TTAAAGTTAT	TCAGTCTTGG	CTGGACACAG	TGCCTCATGT	CTGAAATCCC	2769
AGCACTTTGG	GAGGCTGGGG	CAGGAGGACC	GCTTGAGCCC	CGGGGGTTTG	AAACTGCAGT	2829
GAGCTATGAT	CATGCCACTG	CACTCCAGCC	TAGGTGGCAG	AACTAGACCC	TGTCTCTAAA	2889
AAAACTATTA	TTAGGCCGCG	TGCGGTGGCT	CACGCCTGTA	ATCCCAGCAC	TTTGGGAGAC	2949
TGAGGTGGGT	GGATCACCTG	AGC				2972

(2) INFORMATION FOR SEQ ID NO:115:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 616 amino acids
 - (B) TYPE: amino acid (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: protein
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:115:

Glu Ala Ala His His Leu Val Leu Thr Ala Ser Asp Gly Cly Lys Pro

Pro Arg Ser Ser Thr Val Arg Ile His Val Thr Val Leu Asp Thr Asn

Asp Asn Ala Pro Val Phe Pro His Pro Ile Tyr Arg Val Lys Val Leu

Glu Asn Met Pro Pro Gly Thr Arg Leu Leu Thr Val Thr Ala Ser Asp

Pro Asp Glu Gly Ile Asn Gly Lys Val Ala Tyr Lys Phe Arg Lys Ile 65 70 75 80

Asn Glu Lys Gln Thr Pro Leu Phe Gln Leu Asn Glu Asn Thr Gly Glu Ile Ser Ile Ala Lys Ser Leu Asp Tyr Glu Glu Cys Ser Phe Tyr Glu 100 105 110 Met Glu Ile Gln Ala Glu Asp Val Gly Ala Leu Leu Gly Arg Thr Lys 115 120 125 Leu Leu Ile Ser Val Glu Asp Val Asn Asp Asn Arg Pro Glu Val Ile 130 135 140 Ile Thr Ser Leu Phe Ser Pro Val Leu Glu Asn Ser Leu Pro Gly Thr Val Ile Ala Phe Leu Ser Val His Asp Gln Asp Ser Gly Lys Asn Gly Gln Val Val Cys Tyr Thr Arg Asp Asn Leu Pro Phe Lys Leu Glu Lys Ser Ile Gly Asn Tyr Tyr Arg Leu Val Thr Arg Lys Tyr Leu Asp Arg 195 200 205 Glu Asn Val Ser Ile Tyr Asn Ile Thr Val Met Ala Ser Asp Leu Gly 215 Thr Pro Pro Leu Ser Thr Glu Thr Gln Ile Ala Leu His Val Ala Asp Ile Asn Asp Asn Pro Pro Thr Phe Pro His Ala Ser Tyr Ser Ala Tyr Ile Leu Glu Asn Asn Leu Arg Gly Ala Ser Ile Phe Ser Leu Thr Ala His Asp Pro Asp Ser Gln Glu Asn Ala Gln Val Thr Tyr Ser Val Thr 280 Glu Asp Thr Leu Gln Gly Ala Pro Leu Ser Ser Tyr Ile Ser Ile Asn Ser Asp Thr Gly Val Leu Tyr Ala Leu Gln Ser Phe Asp Tyr Glu Gln Ile Arg Asp Leu Gln Leu Leu Val Thr Ala Ser Asp Ser Gly Asp Pro Pro Leu Ser Ser Asn Met Ser Leu Ser Leu Phe Val Leu Asp Gln Asn Asp Asn Ala Pro Glu Ile Leu Tyr Pro Ala Leu Pro Thr Asp Gly Ser Thr Gly Val Glu Leu Ala Pro Arg Ser Ala Glu Arg Gly Tyr Leu Val Thr Lys Val Val Ala Val Asp Arg Asp Ser Gly Gln Asn Ala Trp Leu 385 390 395 400

Ser Tyr Arg Leu Leu Lys Ala Ser Glu Pro Gly Leu Phe Ser Val Gly 405 410 415

Leu His Thr Gly Glu Val Arg Thr Ala Arg Ala Leu Leu Asp Arg Asp 420 425 430

Ala Leu Lys Gln Ser Leu Val Val Ala Val Gln Asp His Gly Gln Pro 435 440 445

Pro Leu Ser Ala Thr Val Thr Leu Thr Val Ala Val Ala Asp Ser Ile 450 455 460

Pro Glu Val Leu Thr Glu Leu Gly Ser Leu Lys Pro Ser Val Asp Pro 465 470 480

Asn Asp Ser Ser Leu Thr Leu Tyr Leu Val Val Ala Val Ala Ala Ile 485 490 495

Ser Cys Val Phe Leu Ala Phe Val Ala Val Leu Leu Gly Leu Arg Leu 500 505 510

Arg Arg Trp His Lys Ser Arg Leu Leu Gln Asp Ser Gly Gly Arg Leu 515 520 525

Val Gly Val Pro Ala Ser His Phe Val Gly Val Glu Val Gln Ala 530 540

Phe Leu Gln Thr Tyr Ser Gln Glu Val Ser Leu Thr Ala Asp Ser Arg 545 550 555 560

Lys Ser His Leu Ile Phe Pro Gln Pro Asn Tyr Ala Asp Met Leu Ile 565 570 575

Ser Gln Glu Gly Cys Glu Lys Asn Asp Ser Leu Leu Thr Ser Val Asp 580 585 590

Phe His Glu Tyr Lys Asn Glu Ala Asp His Gly Gln Val Ser Leu Val 595 600 605

Leu Cys Leu Leu Leu Ile Ser Arg 610 615